

**Chapter 10. Musculoskeletal and joint diseases**







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**Key**

	<p><b>Red drug</b> see <a href="#">GMMMG RAG list</a>  <i>Click on the symbols to access this list</i></p>
	<p><b>Amber drug</b> see <a href="#">GMMMG RAG list</a>  <i>Click on the symbols to access this list</i></p>
	<p><b>Green drug</b> see <a href="#">GMMMG RAG list</a>  <i>Click on the symbols to access this list</i></p>
	<p><b>If a medicine is unlicensed this should be highlighted in the template as follows</b>  <b>Drug name U</b></p>
	<p><b>Not Recommended</b></p>
	<p><b>Over the Counter</b>          In line with NHS England guidance, GM do not routinely support prescribing for conditions which are self-limiting or amenable to self-care. For further details see <a href="#">GM commissioning statement</a>.</p>
<p><b>Order of Drug Choice</b></p>	<p>Where there is no preferred 1<sup>st</sup> line agent provided, the drug choice appears in alphabetical order.</p>

<b>BNF chapter</b>	<b>10 Musculoskeletal and joint diseases</b>	
<b>Section</b>	<b>10.1 Drugs used in rheumatic diseases and gout</b>	
<b>Subsection</b>	<b>10.1.1 Non-steroidal anti-inflammatory drugs</b>	
<b>First choice</b>	<b>Ibuprofen tablets</b> 200mg; 400mg; 600mg	<a href="#">MHRA DSU: High-dose ibuprofen: small increase in cardiovascular risk, June 2015</a>
<b>Alternatives</b>	<b>Naproxen tablets</b> 250mg; 500mg	
	<b>Diclofenac tablets</b> 50mg	<a href="#">MHRA DSU: diclofenac: new contraindications and warnings, June 2013.</a> <a href="#">MHRA DSU: NSAIDs: cardiovascular risks October 2012.</a> Further evidence that the cardiovascular risk with diclofenac is higher than other non-selective NSAIDs and similar to the selective COX-2 inhibitors
	<b>Etoricoxib tablets</b> 30mg, 60mg, 90mg, 120mg	<a href="#">MHRA DSU: Etoricoxib (Arcoxia): revised dose recommendation for rheumatoid arthritis and ankylosing spondylitis</a> <a href="#">MHRA DSU: etoricoxib: prescribing to patients with high blood pressure, July 2008.</a>
<b>Grey drugs</b> Items which are listed as Grey are deemed not suitable for routine prescribing but may be suitable for a defined patient population	<b>Piroxicam</b> Capsules, orodispersible tablets Systemic piroxicam should be initiated only by specialists as a second-line treatment for arthritis. Patients who currently take piroxicam should be reassessed at a routine appointment.	<b>G<sub>n</sub></b> following specialist initiation <a href="#">Criterion 1 (see RAG list)</a>
	<b>Celecoxib</b> Capsules Only for use in palliative care for the relief of cancer pain.	<b>G<sub>n</sub></b> <a href="#">Criterion 1 (see RAG list)</a>
<b>Do Not Prescribe</b>	<b>Naproxen with esomeprazole</b> Modified-release tablets	<a href="#">Criterion 2 (see RAG list)</a>

**General Guidance applicable to all drugs**

- [NICE CG177: Osteoarthritis: Care and management in adults.](#)
- [NICE NG100: Rheumatoid arthritis in adults: management.](#)
- [CKS \(2013\) NSAID prescribing issues](#)
- [BNF: NSAID-associated ulcers](#)
- [CKS \(2013\): Patients on low dose aspirin](#)
- [MHRA DSU: Non-steroidal anti-inflammatory drugs: reminder on renal failure and impairment, May 2009](#)
- [MHRA DSU: Non-steroidal anti-inflammatory drugs: cardiovascular risk, October 2012](#)
- [MHRA DSU: NSAIDs and coxibs: balancing of cardiovascular and gastrointestinal risks, December 2007](#)

<b>Subsection</b>	<b>10.1.2 Corticosteroids</b>
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**Systemic corticosteroids**

The general actions, uses, and cautions of corticosteroids are described [in the BNF](#).

**Local corticosteroid injections**

<b>First choice</b>	<p><b>Methylprednisolone acetate 40mg/ml</b></p> <p>or</p> <p><b>Methylprednisolone acetate 40mg, lidocaine hydrochloride 10mg/ml</b></p>	
<b>Alternatives</b>	<p><b>Dexamethasone sodium phosphate 3.8mg/ml</b></p>	<p><a href="#">MHRA DSU: Dexamethasone 4 mg/ml injection (Organon Laboratories Limited): reformulation with changes in name, concentration, storage conditions, and presentation, October 2014.</a></p>
	<p><b>Hydrocortisone acetate 25mg/ml</b></p>	
	<p><b>Triamcinolone acetonide 10mg/ml</b> <b>Triamcinolone acetonide 40mg/ml</b></p>	

Subsection	<b>10.1.3 Drugs that suppress the rheumatic disease process</b>	
<b>DMARDS</b>		
<b>First Choice</b>	<b>Methotrexate 2.5mg tablets</b>	<p><b>A</b></p> <p><a href="#">NPSA Alert, June 2006</a>. Improving compliance with oral methotrexate guidelines.</p> <p>When prescribing oral methotrexate only ever prescribe 2.5mg tablets and pay particular attention to correct dosing instructions – <b>weekly</b> dosing.</p> <p>Co-prescribe oral folic acid 5mg weekly with methotrexate (to be taken at least 24 hours after the methotrexate [often 2 -3 days after]) and increase dose if necessary dependant on folate levels.</p>
	<b>Methotrexate 50mg/ml pre-filled Pen injection (Metoject®)</b>	<p><b>R</b> pending homecare arrangements</p> <p><b>Specialist initiation only</b></p> <p>If moving the parenteral formulation to homecare, clinicians should ensure that responsibility for the different aspects of care is clearly defined (in particular blood monitoring).</p> <p><a href="#">MHRA DSU: Methotrexate once-weekly for autoimmune diseases: new measures to reduce risk of fatal overdose due to inadvertent daily instead of weekly dosing, Sept 2020</a></p>
	<b>Sulfasalazine e/c 500mg tablets</b>	<b>A</b>
	<b>Hydroxychloroquine sulphate 200mg tablets</b>	<p><b>A</b></p> <p><a href="#">MHRA DSU: Hydroxychloroquine, chloroquine: increased risk of cardiovascular events when used with macrolide antibiotics; reminder of psychiatric reactions, Feb 2022</a></p>
<b>Alternatives</b>	<b>Leflunomide tablets</b> 10mg; 15mg; 20mg	<p><b>A</b></p> <p><b>Specialist initiation only</b></p>
	<b>Azathioprine tablets</b> 25mg; 50mg	<b>A</b>
	<b>Ciclosporin capsules</b> 10mg; 25mg; 50mg; 100mg	<p><b>A</b></p> <p><a href="#">BNF Online</a> Patients should be stabilised on a particular brand of oral ciclosporin because switching between formulations without close monitoring may lead to clinically important changes in blood-ciclosporin concentration. Prescribing and dispensing of ciclosporin should be by brand name to avoid inadvertent switching.</p>



	<p><b>Mycophenolate U</b> Tablets 500mg, Capsules 250mg</p>	<p><b>A</b></p> <p><a href="#">MHRA DSU: Mycophenolate mofetil, mycophenolic acid: new pregnancy-prevention advice for women and men 2015</a></p> <p><a href="#">MHRA DSU: Mycophenolate mofetil: pure red cell aplasia, July 2009</a></p> <p><a href="#">MHRA DSU: Mycophenolate mofetil (CellCept) and mycophenolic acid: risk of hypogammaglobulinaemia and risk of bronchiectasis, January 2015</a></p>
<p><b>General guidance applicable to all drugs</b></p> <ul style="list-style-type: none"> <li><a href="#">NICE NG100: Rheumatoid arthritis in adults: management.</a></li> </ul>		
<p><b>Biologics (secondary care use only)</b></p>		
<p><b><a href="#">GMMMG High Cost Drugs Pathway for Rheumatoid Arthritis (December 2017)</a></b></p>		
<p><b>First Choices</b> <b>(Follow relevant pathway)</b></p>	<p><b>Abatacept</b> Injection or infusion</p>	<p><b>R</b> Prevents full activation of T-lymphocytes</p> <p><a href="#">NICE TA195: Adalimumab, etanercept, infliximab, rituximab and abatacept for the treatment of rheumatoid arthritis after the failure of a TNF inhibitor.</a></p> <p><a href="#">NICE TA375: Adalimumab, etanercept, infliximab, certolizumab pegol, golimumab, tocilizumab and abatacept for rheumatoid arthritis not previously treated with DMARDs or after conventional DMARDs only have failed</a></p> <p><a href="#">TA715: Adalimumab, etanercept, infliximab and abatacept for treating moderate rheumatoid arthritis after conventional DMARDs have failed</a></p>
	<p><b>Adalimumab</b> First choice: Amgevita® ▼ Alternative: Humira® Solution for injection</p>	<p><b>R</b> TNF inhibitor</p> <p><a href="#">NICE TA195: Adalimumab, etanercept, infliximab, rituximab and abatacept for the treatment of rheumatoid arthritis after the failure of a TNF inhibitor.</a></p> <p><a href="#">TA199: Etanercept, infliximab and adalimumab for the treatment of psoriatic arthritis</a></p> <p><a href="#">NICE TA375: Adalimumab, etanercept, infliximab, certolizumab pegol, golimumab, tocilizumab and abatacept for rheumatoid arthritis not previously treated with DMARDs or after conventional DMARDs only have failed</a></p> <p><a href="#">TA383: TNF-alpha inhibitors for ankylosing spondylitis and non-radiographic axial spondyloarthritis</a></p> <p><a href="#">TA392: Adalimumab for treating moderate to severe hidradenitis suppurativa</a></p> <p><a href="#">TA715: Adalimumab, etanercept, infliximab and abatacept for treating moderate rheumatoid arthritis after conventional DMARDs have failed</a></p>

	<p><b>Apremilast</b> (Otezla®) Tablets</p>	<p><b>R</b> PDE4 Inhibitor <a href="#">TA433: Apremilast for treating active psoriatic arthritis</a> <a href="#">MHRA: Apremilast (Otezla ▼): risk of suicidal thoughts and behaviour</a></p>
	<p><b>Baricitinib ▼</b> (Olumiant®) 2mg and 4mg tablets</p>	<p><b>R</b> JAK inhibitor <a href="#">NICE TA466: Baricitinib for moderate to severe rheumatoid arthritis</a> <a href="#">NICE NG191: COVID-19 rapid guideline: managing COVID-19</a> <a href="#">MHRA DSU: Baricitinib (Olumiant ▼): risk of venous thromboembolism, March 2020</a> <a href="#">MHRA DSU: Baricitinib (Olumiant ▼): increased risk of diverticulitis, particularly in patients with risk factors, Aug 2020</a></p>
	<p><b>Belimumab ▼</b> Infusion</p>	<p><b>R</b> Inhibits activity of B-lymphocyte stimulator <a href="#">TA752: Belimumab for treating active autoantibody-positive systemic lupus erythematosus</a></p>
	<p><b>Certolizumab pegol</b> Solution for injection</p>	<p><b>R</b> TNF inhibitor <a href="#">TA383: TNF-alpha inhibitors for ankylosing spondylitis and non-radiographic axial spondyloarthritis</a> <a href="#">NICE TA415: Certolizumab pegol for treating rheumatoid arthritis after inadequate response to a TNF-alpha inhibitor</a> <a href="#">NICE TA445: Certolizumab pegol and secukinumab for treating active psoriatic arthritis after inadequate response to DMARDs</a></p>
	<p><b>Etanercept</b> Solution for injection</p>	<p><b>R</b> TNF inhibitor <a href="#">NICE TA35: Guidance on the use of etanercept for the treatment of juvenile idiopathic arthritis.</a> <a href="#">NICE TA199: Etanercept, infliximab and adalimumab for the treatment of psoriatic arthritis.</a> <a href="#">TA383: TNF-alpha inhibitors for ankylosing spondylitis and non-radiographic axial spondyloarthritis</a> <a href="#">TA715: Adalimumab, etanercept, infliximab and abatacept for treating moderate rheumatoid arthritis after conventional DMARDs have failed</a></p>
	<p><b>Filgotinib ▼</b> 100mg and 200 mg tablets</p>	<p><b>R</b> JAK inhibitor <a href="#">NICE TA676: Filgotinib for treating moderate to severe rheumatoid arthritis</a></p>

	<p><b>Golimumab</b> Solution for injection</p>	<p><b>R</b> TNF inhibitor <a href="#">NICE TA220: Golimumab for the treatment of psoriatic arthritis.</a> <a href="#">NICE TA225: Golimumab for the treatment of rheumatoid arthritis after the failure of previous disease-modifying anti-rheumatic drugs.</a> <a href="#">TA383: TNF-alpha inhibitors for ankylosing spondylitis and non-radiographic axial spondyloarthritis</a> <a href="#">NICE TA497: Golimumab for treating non-radiographic axial spondyloarthritis</a></p>
	<p><b>Guselkumab</b> Solution for injection</p>	<p><b>R</b> IL-23 inhibitor <a href="#">NICE TA711: Guselkumab for treating active psoriatic arthritis after inadequate response to DMARDs</a> <a href="#">NICE TA815: Guselkumab for treating active psoriatic arthritis after inadequate response to DMARDs</a></p>
	<p><b>Infliximab</b> Infusion</p>	<p><b>R</b> TNF inhibitor <a href="#">NICE TA195: Adalimumab, etanercept, infliximab, rituximab and abatacept for the treatment of rheumatoid arthritis after the failure of a TNF inhibitor.</a> <a href="#">TA199: Etanercept, infliximab and adalimumab for the treatment of psoriatic arthritis</a> <a href="#">NICE TA375: Adalimumab, etanercept, infliximab, certolizumab pegol, golimumab, tocilizumab and abatacept for rheumatoid arthritis not previously treated with DMARDs or after conventional DMARDs only have failed</a> <a href="#">TA383: TNF-alpha inhibitors for ankylosing spondylitis and non-radiographic axial spondyloarthritis</a> <a href="#">TA715: Adalimumab, etanercept, infliximab and abatacept for treating moderate rheumatoid arthritis after conventional DMARDs have failed</a></p>
	<p><b>Ixekizumab</b> Solution for injection</p>	<p><b>R</b> Interleukin 17 antagonist <a href="#">NICE TA537: Ixekizumab for treating active psoriatic arthritis after inadequate response to DMARDs</a> <a href="#">NICE TA718: Ixekizumab for treating axial spondyloarthritis</a></p>
	<p><b>Risankizumab ▼</b> Solution for injection</p>	<p><b>R</b> Interleukin 23 antagonist <a href="#">NICE TA803: Risankizumab for treating active psoriatic arthritis after inadequate response to DMARDs</a></p>
	<p><b>Rituximab</b> Infusion</p>	<p><b>R</b> Anti-lymphocytic - causes lysis of B lymphocytes <a href="#">MHRA DSU: Rituximab: progressive multifocal leukoencephalopathy in a patient without prior treatment for rheumatoid arthritis, December 2009.</a></p>

		<p><a href="#">MHRA DSU: Rituximab: screen for hepatitis B virus before treatment, December 2013</a></p> <p><a href="#">NICE TA308: Rituximab in combination with glucocorticoids for treating anti-neutrophil cytoplasmic antibody-associated vasculitis.</a></p> <p><a href="#">GMMMG Rituximab biosimilar recommendation (August 2017)</a></p>
	<p><b>Sarilumab ▼</b> Solution for injection</p>	<p><b>R</b> Interleukin-6 antagonist</p> <p><a href="#">NICE TA485: Sarilumab for moderate to severe rheumatoid arthritis</a></p> <p><a href="#">NG191: managing COVID-19</a></p>
	<p><b>Secukinumab</b> Solution for injection</p>	<p><b>R</b></p> <p><a href="#">NICE TA407: Secukinumab for active ankylosing spondylitis after treatment with non-steroidal anti-inflammatory drugs or TNF-alpha inhibitors</a></p> <p><a href="#">NICE TA719: Secukinumab for treating non-radiographic axial spondyloarthritis</a></p>
	<p><b>Tocilizumab</b> Injection, infusion</p>	<p><b>R</b> Interleukin-6 antagonist</p> <p><a href="#">NICE TA238: Systemic juvenile idiopathic arthritis – tocilizumab.</a></p> <p><a href="#">NICE TA247: Tocilizumab for the treatment of systemic juvenile idiopathic arthritis.</a></p> <p><a href="#">NICE TA518: Tocilizumab for treating giant cell arteritis</a></p> <p><a href="#">NG191: managing COVID-19</a></p>
	<p><b>Tofacitinib ▼</b> 5mg and 10mg tablets</p>	<p><b>R</b> JAK inhibitor</p> <p><a href="#">NICE TA480: Tofacitinib for moderate to severe rheumatoid arthritis</a></p> <p><a href="#">NICE TA543: Tofacitinib for treating active psoriatic arthritis after inadequate response to DMARDs</a></p> <p><a href="#">MHRA DSU: Tofacitinib (Xeljanz ▼): new measures to minimise risk of venous thromboembolism and of serious and fatal infections, March 2020</a></p> <p><a href="#">MHRA DSU: Tofacitinib (Xeljanz ▼): new measures to minimise risk of major adverse cardiovascular events and malignancies</a></p>
	<p><b>Upadacitinib ▼</b> 15 mg prolonged-release tablets</p>	<p><b>R</b> JAK inhibitor</p> <p><a href="#">NICE TA665: Upadacitinib for treating severe rheumatoid arthritis</a></p> <p><a href="#">NICE TA744: Upadacitinib for treating moderate rheumatoid arthritis</a></p> <p><a href="#">NICE TA768: for treating active psoriatic arthritis after inadequate response to DMARDs</a></p>



	<b>Ustekinumab</b> Solution for injection	 <a href="#">NICE TA340: Ustekinumab for treating active psoriatic arthritis.</a>
<b>General guidance applicable to all drugs</b> <ul style="list-style-type: none"> <li>• Biologics in Rheumatoid Arthritis can be used, without prior funding approval, according to <a href="#">GMMMG approved pathways</a>: High Cost Drugs Pathway for Rheumatoid Arthritis and Harmonised Biologics Pathway for Ankylosing Spondylitis and Psoriatic Arthritis</li> <li>• <a href="#">MHRA DSU</a>: Tumour necrosis factor alpha inhibitors: risk of tuberculosis—screen all patients before starting treatment and monitor them closely, April 2014</li> <li>• <a href="#">MHRA DSU: TNFa inhibitors: risk of TB – screen all patients before starting treatment and monitor them closely (April 2016)</a></li> </ul>		
<b>Additional NICE guidance and technology appraisals:</b> <ul style="list-style-type: none"> <li>• <a href="#">NICE NG100: Rheumatoid arthritis in adults: management.</a></li> <li>• <a href="#">NICE TA383: TNF-alpha inhibitors for ankylosing spondylitis and non-radiographic axial spondyloarthritis</a></li> </ul>		
<b>Subsection</b>	<b>10.1.4 Gout and cytotoxic induced hyperuricaemia</b>	
<b>General guidance applicable to all drugs</b> <ul style="list-style-type: none"> <li>• <a href="#">NICE NG219: Gout: diagnosis and management</a></li> <li>• <a href="#">British Society for Rheumatology Guideline for the Management of Gout</a></li> </ul>		
<b>NICE Technology Appraisals:</b> <ul style="list-style-type: none"> <li>• <a href="#">NICE TA291: Pegloticase for treating severe debilitating chronic tophaceous gout</a></li> </ul>		
<b>Acute attack</b>		
<b>First choice</b> Alternative where NSAIDS contraindicated	<b>Colchicine 500microgram tablets</b>	<a href="#">MHRA DSU: colchicine: extremely toxic in overdose, November 2009.</a>
<b>Long-term control of gout</b>		
<b>First choice</b>	<b>Allopurinol tablets</b> 100mg; 300mg	
<b>Alternatives</b>	<b>Febuxostat tablets</b> 80mg; 120mg	 Use only where allopurinol is contraindicated or not tolerated. (As per NICE guidance and/or on specialist advice.) <a href="#">MHRA DSU: Febuxostat (Adenuric▼): stop treatment if signs or symptoms of serious hypersensitivity occur, June 2012.</a>
	<b>Sulfinpyrazone tablets</b> 100mg; 200mg	
<b>Do Not Prescribe</b>	<b>Lesinurad</b>	<a href="#">Criterion 2 (see RAG list)</a>

	Tablets	
<b>Subsection</b>	<b>10.1.5 Other drugs for rheumatic diseases</b>	
<p><b>Applicable guidance:</b></p> <ul style="list-style-type: none"> <li>• <a href="#">NICE CG177: Osteoarthritis: Care and management in adults</a>. - section 1.4.5 Do not offer glucosamine or chondroitin products for the management of osteoarthritis.</li> <li>• GMMMG 'Do Not Prescribe and Grey Lists': <a href="#">glucosamine</a> and <a href="#">synovial fluid injections</a> (including hyaluronan and sodium hyaluronate injection).</li> <li>• <a href="#">GM Policy Statement (2020). Hyaluronic Acid injections for Osteoarthritis. (GM037)</a></li> </ul> <p><b>NICE Technology Appraisals:</b></p> <ul style="list-style-type: none"> <li>• <a href="#">NICE TA477: Autologous chondrocyte implantation for treating symptomatic articular cartilage defects of the knee</a></li> <li>• <a href="#">NICE TA508: Autologous chondrocyte implantation using chondrosphere for treating symptomatic articular cartilage defects of the knee</a></li> </ul>		
<b>Do Not Prescribe</b>	<p><b>Glucosamine</b> With or without chondroitin</p>	<a href="#">Criterion 1 (see RAG list)</a>
	<p><b>Synovial fluid injections</b> Including hyaluronan and sodium hyaluronate injections</p>	<a href="#">Criterion 1 (see RAG list)</a>

<b>Section</b>	<b>10.2 Drugs used in neuromuscular disorders</b>	
<b>Subsection</b>	<b>10.2.1 Drugs that enhance neuromuscular transmission</b>	
<b>First choice</b>	<b>Pyridostigmine</b> tablets: 60mg	
<b>Alternatives</b>	<b>Neostigmine</b> tablets: 15mg	
<b>Subsection</b>	<b>10.2.2 Skeletal muscle relaxants</b>	
<b>First choice</b>	<b>Dantrolene</b> capsules: 25mg, 100mg	
<b>Alternatives</b>	<b>Diazepam</b> tablets 2mg, 5mg, 10mg oral solution 2mg/5ml	
	<b>Baclofen</b> tablets: 10mg oral solution: 5mg/5ml	
	<b>Mexiletine</b> capsules: 167mg	<b>R</b> <a href="#">NICE TA748: Mexiletine for treating the symptoms of myotonia in non-dystrophic myotonic disorders</a>
	<b>Tizanidine</b> tablets: 2mg, 4mg	<b>G<sub>n</sub></b> Following specialist initiation
<b>Section</b>	<b>Miscellaneous</b>	
	<b>Nusinersen</b> (Prevymis®) tablets: 240mg	<b>R</b> <a href="#">NICE TA588: Nusinersen for treating spinal muscular atrophy</a>
	<b>Risdiplam</b> (Evrysdi®▼) powder for oral solution 0.75mg/ml	<b>R</b> <a href="#">NICE TA755: Risdiplam for treating spinal muscular atrophy</a>
<b>Do Not Prescribe</b>	<b>Therabite®</b> Jaw rehabilitation system	<a href="#">Criterion 1 (see RAG list)</a>

<b>Section</b>	<b>10.3 Drugs for the relief of soft-tissue inflammation</b>	
<b>Subsection</b>	<b>10.3.1 Enzymes</b> (Secondary care use)	
	<b>Collagenase (Xiapex®)</b>	<b>R</b> <a href="#">NICE TA459: Collagenase clostridium histolyticum for treating Dupuytren's contracture</a>
<b>Subsection</b>	<b>10.3.2 Rubefaciants and other topical antirheumatics</b>	
<b>Rubefaciants</b>		
<b>Do Not Prescribe</b>	<b>Rubefaciants</b> Topical rubefacient products may contain nicotinate and salicylate compounds, essential oils, capsicum, and camphor. Topical NSAID or capsaicin preps are not rubefaciants	<b>Criterion 1 (see <a href="#">RAG list</a>)</b>
<b>Topical NSAIDS</b>		
<b>Options</b>	<b>Ibuprofen 5% Gel</b> (Available as 30g, 50g or 100g tubes)	
	<b>Ketoprofen 2.5% Gel</b> (Available as 30g, 50g or 100g tubes)	<a href="#">MHRA DSU: topical ketoprofen: reminder on photosensitivity reactions, June 2009</a> and <a href="#">MHRA August 2010:</a>
	<b>Piroxicam 0.5% Gel</b> (Available as 60g or 112g tubes)	
<b>Capsaicin</b>		
<b>Options</b>	<b>Capsaicin 0.025% cream</b>	<a href="#">NICE CG177: Osteoarthritis: Care and management in adults.</a>  <a href="#">GMMMG Neuropathic Pain Guideline November 2019</a>
	<b>Capsaicin 0.075% cream</b>	
<b>Poultices</b>		
Not recommended for prescribing		