(Collaboration Commissioners' list of Indications that may be commissioned for excluded drugs 2019-20)

Version 14.10 CCG National Tariff Excluded Drugs (NTeD) List

Provisional list subject to publication of High Cost Drug exclusion list and and Indications for NHS England drugs list (Medicines not reimbursed through national prices and directly commissioned by NHS England) High cost drugs excluded from the National Tariff are not commissioned by NCL Clinical Commissioning Groups (CCGs) unless the drugs and indications have approval through local processes.

Please note that the enclosed information reflects the current commissioning arrangements. These arrangements are subject to change in year if commissioning responsibility for particular drugs or services change between NCL CCGs and the NHSE and / or new local drug policies are introduced or new NICE guidance is

The pathways refered to in this document are avaiable on the JFC website accessible here: https://www.ncl-mon.nhs.uk/documentation/moc/prescribing-policies/

The tables in Appendix 1 summarise the policy on the commissioning of drugs. It is coded by a "traffic light" scheme as follows:

| | Not comm | issioned by CCG. When | e provider Trusts prescribe this treatment the funding res | ponsibility lies with th | ne provider and not the | commissioner. | |
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| | IFR appro | | | | | | |
| | | | | | | | or approval required using proforma or electronic Blueteq® forms as per contract. Clinical evidence of outcomes may |
| | | | | | | | er minimum dataset information must be provided on back up data to invoices. |
| | | | ty. CCGs should not consider funding requests for these on missioning/spec-services/key-docs/ | drugs in these indicat | ions / group of patients | . For drugs or indications wh | ere NHS England is the responsible commissioner the current version of the NHS England drug list should be refered |
| | nttp://ww | w.engiand.nns.uk/comr | hissioning/spec-services/key-docs/ | | | | |
| Drug name | BNF section | Category | Licensed Indication/ Indication | NICE Guidance | Indications agreed for funding across | Is this commissioned? What is required for | Notes |
| Abaloparatide | 6.4 | Analogue of human parathyroid hormone- related protein | Osteoporosis- post-menopausal, with high risk of fracture | ID882 | NOT COMMISSIONED | funding approval? NOT COMMISSIONED | Not yet launched in U.K. |
| Abatacept | 10.1 | Cytokine modulators | Rheumatoid arthritis after failure of anti-TNF | NICE TA195, Aug 2010 | In line with NICE TA195 criteria or NC RA pathway. | Complete TBF/blueteq (using agreed local systems) | Adults only. GP prescribing NOT expected. NHS England is responsible commissioner for paediatric use until patient are transferred into adult services. |
| Abatacept | 10.1 | Cytokine modulators | Juvenile arthritis | NICE TA373, Dec 2015. | | NOT COMMISSIONED | NHS England is responsible commissioner for paediatric use until patients are transferred into the adult service. |
| Abatacept | 10.1 | Cytokine modulators | Rheumatoid arthritis not previously treated with DMARDs or after conventional DMARDs only have failed | NICE TA375, Jan 2016 | In line with NICE TA375 criteria and NCL RA pathway. | Complete TBF/blueteq (using agreed local systems) | Adults only. GP prescribing NOT expected. NHS England is responsible commissioner for paediatric use. |
| Abatacept | 10.1 | Cytokine modulators | Psoriatic arthritis | No NICE guidance anticipated. | NOT COMMISSIONED | NOT COMMISSIONED | Adults only. GP prescribing NOT expected. NHS England is responsible commissioner for paediatric use. |
| Abatacept | 10.1 | Cytokine modulators | All other indications | No NICE guidance anticipated. | NOT COMMISSIONED | NOT COMMISSIONED | NHS England is responsible commissioner for paediatric use. |
| Adalimumab | 10.1 | Cytokine modulators | Rheumatoid arthritis | NICE TA375, Jan 2016 or TA195, Aug 2010. | In line with NICE TA375 or TA195 criteria or NCL RA pathway | Complete TBF/blueteq (using agreed local systems) | Adults only. GP prescribing NOT expected. NHS England is responsible commissioner for paediatric use until patient are transferred into adult services, Uveitis and Hidradenitis Suppurativa |
| Adalimumab | 10.1 | Cytokine modulators | Psoriatic Arthritis | NICE TA199, Aug 2010 | In line with NICE criteria TA199 or NCL Psoriasis pathway (awaited as per JFC) | Complete TBF/blueteq (using agreed local systems) | Adults only. GP prescribing NOT expected. NHS England is responsible commissioner for paediatric use, uveitis and Hidradenitis Suppurativa |
| Adalimumab | 10.1 | Cytokine modulators | Ankylosing spondylitis | NICE TA383, Feb 2016. | In line with NICE TA383 criteria. | Complete TBF/blueteq (using agreed local systems) | Adults only. GP prescribing NOT expected. NHS England is responsible commissioner for paediatric use, uveitis and Hidradenitis Suppurativa |
| Adalimumab | 13.3.1 | Cytokine modulators | Psoriasis | NICE TA146, June 2008 | In line with NICE TA146 criteria or NCL Psoriasis pathway (awaited as per JFC) | Complete TBF/blueteq (using agreed local systems) | Adults only. GP prescribing NOT expected. NHS England is responsible commissioner for paediatric use, uveitis and Hidradenitis Suppurativa |
| Adalimumab | 1.1.3 | Cytokine modulators | Crohn's Disease | NICE TA187, May 2010 | In line with NICE TA187 criteria. | Complete TBF/blueteq (using agreed local systems) | Adults only. GP prescribing NOT expected. NHS England is responsible commissioner for paediatric use, uveitis and Hidradenitis Suppurativa |
| dalimumab | 10.1 | Cytokine modulators | Axial spondylo-arthritis (non-radiographic) | NICE TA383, Feb 2016. | In line with NICE TA383 criteria. | Complete TBF/blueteq (using agreed local systems) | Adults only. GP prescribing NOT expected. NHS England is responsible commissioner for paediatric use, uveitis and Hidradenitis Suppurativa |
| dalimumab | 1.1.3 | Cytokine modulators | Moderately to severely active ulcerative colitis after the failure of conventional therapy | NICE TA329, Feb 2015 | In line with NICE TA329 criteria. | Complete TBF/blueteq (using agreed local systems) | Adults only. GP prescribing NOT expected. NHS England is responsible commissioner for paediatric use, uveitis and Hidradenitis Suppurativa |
| Adalimumab | 1.1.3 10.1 13.3.1 | Cytokine modulators | All other indications. | NICE TA460, July 2017 NICE TA392, June 2016 NICE TA455, July 2017 NICE TA 373, DEC 2015 | COMMISSIONED | NOT COMMISSIONED | NHS England is responsible commissioner for paediatric use where an adult TA is available and for JIA until patients are transferred to adult services. NHS England is responsible commissioner for uveitis in adults, bechet's, hidradenit suppurativa and. |

| Recognition of the company of the | Drug name | BNF | Category | Licensed Indication/ Indication | NICE Guidance | Indications agreed | Is this commissioned? | Notes |
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| migraine migraines 2012 TA260 criteria. (using agreed local | Bortezomio Botulinum Toxin A | 4.6.1 | Prophylaxis of | | NICE TA260, June | In line with NICE | | |
| | | | | | | | (using agreed local | |
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| Drug name | BNF section | Category | Licensed Indication/ Indication | NICE Guidance | Indications agreed for funding across NCL | Is this commissioned? What is required for funding approval? | Notes |
|---|----------------|--|---|--|--|--|---|
| Botulinum Toxin A | 4.1 | Essential tremor, chorea, tics and related disorders | Laryngeal dystonia (spasmodic dystonia), hydradenitis suppurative, mechanical neck disorders, correction of strabismus (squint) in paediatrics, dysphagia (causes other than achalasia), Raynaud's Disease, Massateric Hypertrophy. | No NICE guidance anticipated | NOT COMMISSIONED | NOT COMMISSIONED. | Historical arrangement for BCF to reimburse for the following additional indication: achalasia, anal fissures, axillary hyperhidrosis, cerebral palsy, cervical dystonia, foream dystonia, hemifacial spasm, severe blepharospasm |
| Botulinum Toxin A | 4.1 | Essential tremor, chorea, tics and related disorders | All other indications | No NICE guidance anticipated | NOT COMMISSIONED | NOT COMMISSIONED | Historical arrangement for BCF to reimburse for the following additional indication: achalasia, anal fissures, axillary hyperhidrosis, cerebral palsy, cervical dystonia, foream dystonia, hemifacial spasm, severe blepharospasm |
| Botulinum Toxin A | 4.1 | Essential tremor, chorea, tics and related disorders | Chronic anal fissure, severe blepharospasm, hemifacial spasm, cervical dystonia (spasmodic torticollis), focal spasticity in upper and lower limb in adults (causes other than stroke), hyperhidrosis of the axillae, dysphagia caused by achalasia, overactive bladder, spasticity treatment in paediatric cerebral palsy, hypersalivation caused by disease and Hirschsprung's disease. | No NICE guidance anticipated.NICE TA ID768 no due date. | For RF & UCLH, indications covered under voice are Spasmodic /spatic dysphonia, tourette's disorder affecting the larynx, laryngeal granuloma refractory to surgery. Also spasticity and dystonia associated with human prion disease, stroke, trauma /spinal injury. (JFC Jan 2013) | Brand and Indication must be provided in SLAM | GP prescribing NOT expected.NHS England is responsible commissioner for focal spasticity in children. Treatment of spasticity in paediatric cerebral palsy is commissioned by NHSE at specialist centres. NHS England is responsible commissioner for intravesical use in spinal cord injury. |
| Botulinum Toxin B | 4.1 | Torsion, dystonias and other involuntary movements | Spasmodic torticollis (Cervical dystonia) | | | | |
| Brodalumab | 13.3.1 | Drugs affecting the immune response | Moderate to severe plaque psoriasis | NICE TA511, Mar 2018 | In line with NICE TA511 criteria. | Complete Blueteq form (using agreed local systems) | Adults only. GP prescribing NOT expected. NHS England is responsible commissioner for paediatric use through the IFR process. |
| Brodalumab | 10.1 | Drugs affecting the immune response | Psoriatic arthritis | No NICE guidance anticipated. | NOT COMMISSIONED | NOT COMMISSIONED | |
| Canakinumab | 8.1 | Immunomodulating Drugs | Cryopyrin Associated Periodic Syndroms | No NICE guidance anticipated. | NOT COMMISSIONED | NOT COMMISSIONED | NHS England responsible commissioner as per service specification (highly specialised criteria only). |
| Canakinumab | 10.1 | Immunomodulating Drugs | Juvenile Arthritis | No NICE guidance anticipated. | NOT ROUTINELY COMMISSIONED | NOT ROUTINELY COMMISSIONED | NHS England responsible commissioner for paediatric use through the IFR process. |
| Certolizumab Pegol | 10.1 | Cytokine modulators | Rheumatoid arthritis not previously treated with DMARDs or after conventional DMARDs only have failed | NICE TA375, Jan 2016 | In line with NICE TA375 criteria. | Complete TBF/Blueteq (using agreed local systems) | Adults only, GP prescribing NOT expected. NHS England responsible commissioner for paediatric use through the IFR process. |
| Certolizumab Pegol | 10.1 | Cytokine modulators | Rheumatoid arthritis after TNF-alpha inhibitor | NICE TA415, Oct 2016 | in line with NICE TA415 criteria. | Complete TBF/Blueteq (using agreed local systems) | Adults only, GP prescribing NOT expected. NHS England responsible commissioner for paediatric use through the IFR process. |
| Certolizumab Pegol | 10.1 | Cytokine modulators | Active psoriatic arthritis after DMARDs | NICE TA445, May 2017 | In line with NICE TA445 criteria. | Complete TBF/Blueteq (using agreed local systems) | Adults only, GP prescribing NOT expected. NHS England responsible commissioner for paediatric use through the IFR process. |
| Certolizumab Pegol | 10.1 | Cytokine modulators | Ankylosing spondylitis and non-radiographic axial spondyloarthritis | NICE TA383, Feb 2016 | In line with NICE TA383 criteria. | Complete TBF/Blueteq (using agreed local systems) | Adults only, GP prescribing NOT expected. NHS England responsible commissioner for paediatric use through the IFR process. |
| Certolizumab Pegol | 10.1 | Cytokine modulators | All other indications. | No NICE guidance anticipated. | No NICE guidance anticipated. | NOT COMMISSIONED | |
| Certolizumab Pegol | 13.3.1 | Cytokine modulators | Moderate to severe plaque psoriasis | NICE ID1232, expected April 2019 | NOT COMMISSIONED | NOT COMMISSIONED | |
| Co- careldopa intestinal gel (Duodopa®) | 4.4.2 | Parkinson's disease | Management of advanced Parkinson's disease | NHS England | NHS ENGLAND CLINICAL COMMISSIONING POLICY: D04/P/e | NHS England | NHS England is the responsible commissioner. |
| Colistimethate sodium dry powder for inhalation (Colobreathe®) | 5.2 | Antibacterial drugs | Cystic Fibrosis | NICE TA276, March 2013. | NHSE Policy -Inhaled Therapy For Adults And Children With Cystic Fibrosis Policy Ref: A01/P/b | NHS England | Only excluded when given by nebulisation/inhalation. No GP prescribing of Colobreathe® as Patient Access Scheme not available in primary care. |
| Colistimethate sodium powder (Promixin®) /injection for nebulisation | 5.1.7 | Antibacterial drugs | Cystic Fibrosis | NICE TA276, March 2013. | NHSE Policy -Inhaled Therapy For Adults And Children With Cystic Fibrosis Policy Ref: A01/P/b | NHS England | Only excluded when given by nebulisation/inhalation. |

| Drug name | BNF section | Category | Licensed Indication/ Indication | NICE Guidance | Indications agreed for funding across NCL | Is this commissioned? What is required for funding approval? | Notes |
|--|----------------|------------------------------------|---|---|--|--|--|
| Collagenase | 10.5.2 | Enzymes | Dupuytren's Contracture | NICE TA459, July 2017 | In line with NICE TA459 criteria. | (using agreed local systems) | Only excluded from tariff when used in outpatient |
| Collagenase | 10.5.2 | Enzymes | Peyronie's disease (PsD) | N/A | NOT COMMISSIONED | NOT COMMISSIONED | |
| Deferasirox | 9.2 | Iron overload | Anaemia related to chronic iron overload for myelodysplastic syndrome only | No NICE guidance anticipated. | Local policy being updated for MDS only | Complete proforma (using agreed local systems) | NHS England responsible commissioner for thalassemia and sickle cell disease |
| Deferiprone | 9.2 | Iron overload | Anaemia related to chronic iron overload for myelodysplastic syndrome only | No NICE guidance anticipated. | Local policy being updated for MDS only | Complete proforma (using agreed local systems) | NHS England responsible commissioner for thalassemia and sickle cell disease |
| Desferrioxamine | 9.2 | Iron overload | Anaemia related to chronic iron overload for myelodysplastic syndrome only | NICE TA suspended | Local policy being updated for MDS only | agreed local systems) | NHS England responsible commissioner for thalassemia and sickle cell disease. |
| Dexamethasone intravitreal implant | 11.6.2 | Macular oedema | | NICE TA229, July 2011. | TA229 criteria. | Complete TBF/Blueteq (using agreed local systems) | |
| Dexamethasone intravitreal implant | 11.6.2 | Macular oedema | Diabetic Macular Oedema | NICE TA349, July 2015 | In line with NICE TA349 criteria or DMO pathway. | Complete TBF/Blueteq (using agreed local systems) | |
| Dexamethasone intravitreal implant | 11.6.2 | Macular oedema | Inflammation of the posterior segment of the eye presenting as non-infectious uveitis | NICE TA460, July 2017 | In line with NICE TA460 criteria. | Complete TBF/Blueteq (using agreed local systems) | |
| Dibotermin alfa | N/A | Bone Morphogenetic Protein(BMP) | Acute tibia fractures | No NICE guidance anticipated. | BMP for secondary revision of spinal fusion surgery as per local commissioning policy letter dated Dec 2013 | Complete proforma (using agreed local systems) | NHS England responsible commissioner for complex spinal surgery. |
| Digoxin immune fab | 16.3.2 | Poisoning | Life - threatening digoxin toxicity | National Poisons Centre Guidelines | Only commissioned in line with National Poison Centres guidelines. | Notification on SLAM. | Available on named patient basis only in the hospital. |
| Dimethyl fumarate | 13.3.1 | Preparations for psoriasis | Chronic plaque psoriasis, moderate-to-severe, modified- release prep | NICE TA475, Sept 2017 | | Complete Blueteq form (using agreed local systems) | |
| Dimethyl fumarate | 13.3.1 | Preparations for psoriasis | Plaque psoriasis, moderate to severe | NICE TA475, Sept 2017 | In line with NICE TA475 criteria. | Complete TBF/Blueteq (using agreed local systems) | |
| Dimethyl fumarate | 8.1.1 | Immunomodulating Drugs | relapsing-remitting multiple sclerosis | NICE TA320, Aug 2014 | In line with NICE TA320 criteria. | NHS England | NHS England is responsible for commissioning for MS. |
| Dornase alpha | 3.3.1 | Mucolytics | Cystic fibrosis | NHS England | In line with NHS England Policy: A01/P/b. | NHS England | NHS England is the responsible commissioner for cystic fibrosis. NHS England is also responsible for primary ciliary dyskinesia through IFR process. |
| Dupilumab | 13.3.1 | Cytokine modulators | Moderate to severe atopic dermatitis in adults | NICE TA534, Aug 2018 | In line with NICE TA534 criteria. | Complete TBF/blueteq (using agreed local systems) | NHS England is the responsible commissioner for use in astham through the IFR process. |
| Eculizumab | 9.1.1a | Immunosuppressants | Paroxysmal noctumal Haemoglobinuria | NHS England | In line with NHS England specialised commissioning team service specification | NHS England | NHS England is the commissioner for indications included within this category. The indications also include: - Organ rejection post kidney transplant (policy A07/P/b): Not routinely commissioned, through NHSE IFR process C3 Glomerulopathy (post transplant) (clinical commissioning policy16054/P) - AHUS (clinical commissioning policy E03/PS(HSS)/A NICE HST1) |
| Eltrombopag | 9.4.2 | Platelet disorder drugs | Chronic idiopathic thrombocytopenic purpura (ITP) | NICE TA293, July 2013 | In line with NICE TA293 criteria. | Complete TBF/blueteq (using agreed local systems) | GP prescribing NOT expected. NHS England is the responsible commissioner for paediatric |
| Eltrombopag | 9.4.2 | Platelet disorder drugs | Thrombocytopenia in adult patients with chronic hepatitis C virus infection | No NICE guidance anticipated. | NOT COMMISSIONED | NOT COMMISSIONED | NHS England is responsible commissioner for paediatric use |
| Eltrombopag | 9.4.2 | Platelet disorder drugs | Acquired severe aplastic anaemia (SAA) refractory to prior immunosuppressive therapy | NICE TA terminated | NOT COMMISSIONED. | NOT COMMISSIONED. | NICE was unable to make recommendations on eltrombopag for severe aplastic anaemia refractory to immunosuppressive therapy because no evidence submission was received. |
| Eptotermin alfa | N/A | Bone morphogenic protein | Complex spinal surgery | N/A | NOT ROUTINELY COMMISSIONED | NOT ROUTINELY COMMISSIONED | NHS England is the responsible commissioner through the IFR process. |
| Eptotermin alfa | N/A | Bone morphogenic | Acute tibia fracture (adults) | NOT COMMISSIONED | NOT COMMISSIONED | NOT COMMISSIONED | |
| Erythropoeitin for non-dialysis renal patients | 9.1.1 | Drugs used in renal anaemias | Pre-dialysis | No NICE guidance anticipated. | NOT COMMISSIONED | NOT COMMISSIONED | |
| Epoetin (all variants) | 9.1.1 | Drugs used in renal anaemias | Dialysis-induced anaemia | NICE CG114 | Renal Dialysis Only as per NICE CG114. | NHS England | NHS England is the responsible commissioner for dialysis-induced anaemia. |
| Etanercept or licensed biosimilar | 10.1 | Cytokine modulators | Rheumatoid arthritis | NICE TA375, Jan 2016 or TA195 Aug 2010. | In line with NICE TA373 or TA195 criteria or NCL RA pathway. | Complete TBF/Blueteq (using agreed local systems) | NHS England is responsible commissioner for paediatric use, until patients are transferred into the adult service. |

| Drug name | BNF | Category | Licensed Indication/ Indication | NICE Guidance | Indications agreed | | Notes |
|-----------------------------------|---------|---|---|---|---|---|---|
| | section | | | | for funding across NCL | What is required for funding approval? | |
| Etanercept or licensed biosimilar | 10.1 | Cytokine modulators | Ankylosing spondylitis (adults) | NICE TA383, Feb | In line with NICE | Complete TBF/Blueteq | NHS England is responsible commissioner for paediatric use. |
| | | | | 2016. | TA383 criteria. | (using agreed local systems) | |
| Etanercept or licensed biosimilar | 10.1 | | Axial spondyloarthritis (non- radiographic) | NICE TA383, Feb 2016. | TA383 criteria. | Complete TBF/Blueteq (using agreed local systems) | NHS England is responsible commissioner for paediatric use. |
| Etanercept or licensed biosimilar | 10.1 | Cytokine modulators | Psoriatic Arthritis | NICE TA199, Aug 2010 | In line with NICE TA199 criteria. | Complete TBF/Blueteq (using agreed local systems) | NHS England is responsible commissioner for paediatric use. |
| Etanercept or licensed biosimilar | 13.3.1 | Cytokine modulators | Psoriasis | NICE TA103, July 2006 | In line with NICE TA103 criteria or NCL Psoriasis pathway (awaited as per JFC) | Complete TBF/Blueteq (using agreed local systems) | NHS England is responsible commissioner for paediatric use. |
| Etanercept or licensed biosimilar | 10.1 | Cytokine modulators | All other indications | No NICE guidance anticipated. | NOT COMMISSIONED. | NOT COMMISSIONED. | NHS England is responsible commissioner for paediatric use. |
| Evolocumab | 2.6 | PCSK9 Monoclonal Antibody Inhibitor | Primary hypercholesterolaemia or mixed dyslipidaemia (heterozygous familial and non-familial) | NICE TA394, Jun 2016 | In line with NICE TA394 criteria. | Complete TBF/Blueteq (using agreed local systems) | |
| Evolocumab | 2.6 | PCSK9 Monoclonal Antibody Inhibitor | Homozygous familial hypercholesterolemia | 2016 | NHS England | NHS England | NHS England is the responsible commissioner. |
| Fluocinolone acetonide | 11.6.2 | Macular oedema | Chronic diabetic macular oedema | NICE TA301, Nov 2013 | in line with NICE TA301 criteria. | Complete TBF/Blueteq (using agreed local systems) | |
| Fomepizole | N/A | Poisoning | Poisoning | National Poisons Centre Guidelines | Only commissioned in line with National Poison Centres guidelines. | Notification on SLAM. Check with CCGs | Available on named patient basis only in the hospital. |
| Golimumab | 10.1 | Cytokine modulators | Psoriatic Arthritis | NICE TA220, April 2011 | In line with NICE TA220 criteria. | Complete TBF/Blueteq (using agreed local systems) | NHS England is responsible commissioner for paediatric use. |
| Golimumab | 10.1 | Cytokine modulators | Rheumatoid arthritis | NICE TA375, Jan 2016 and TA225, June 2011 | In line with NICE TA375 or TA225 criteria or NCL RA pathway | Complete TBF/Blueteq (using agreed local systems) | NHS England is responsible commissioner for paediatric use. |
| Golimumab | 10.1 | Cytokine modulators | Ankylosing spondylitis | NICE TA383, Feb 2016 | In line NICE TA383 criteria. | Complete TBF/Blueteq (using agreed local systems) | NHS England is responsible commissioner for paediatric use. |
| Golimumab | 1.1.3 | Cytokine modulators | Moderately to severely active ulcerative colitis after the failure of conventional therapy | NICE TA329, Feb 2015 | In line with NICE TA329 criteria. | Complete TBF/Blueteq (using agreed local systems) | NHS England is responsible commissioner for paediatric use. |
| Golimumab | 10.1 | Cytokine modulators | Axial spondyloarthritis (non- radiographic) | NICE TA383, Feb 2016 and NICE TA497, Jan 2018 | In line with NICE TA383 and TA497 criteria. | Complete TBF/Blueteq (using agreed local systems) | NHS England is responsible commissioner for paediatric use. |
| Golimumab | 10.1 | Cytokine modulators | All other indications | No NICE guidance anticipated. | NOT COMMISSIONED. | NOT COMMISSIONED. | NHS England is responsible commissioner for paediatric use. |
| Guselkumab | 13.3.1 | Cytokine modulators | Moderate to severe plaque psoriasis (adults) | NICE TA521, Jun 2018 | In line with NICE TA521 criteria. | Complete TBF/Blueteq (using agreed local systems) | |
| lloprost | 2.4.1c | Vasodilator antihypertensive drugs/Pulmanory Arterial Hypertension | Critical limb ischaemia | No NICE guidance anticipated. | NOT COMMISSIONED. | NOT COMMISSIONED | NHS England is the responsible commissioner for PAH. |
| lloprost | 2.4.1c | Vasodilator antihypertensive drugs/Pulmanory Arterial Hypertension | Pulmonary Arterial Hypertension | NHS England Policy | In line with NHS England clinical commissioning policy: A11/P/c | NHS England | NHS England is the responsible commissioner. |
| Infliximab or licensed biosimilar | 1.1.3 | Cytokine modulators | Crohn's disease | NICE TA187, May 2010 | | Complete TBF/Blueteq (using agreed local systems) | NHS England is responsible commissioner for paediatric use. |
| Infliximab or licensed biosimilar | 10.1 | Cytokine modulators | Rheumatoid arthritis | NICE TA375, Jan 2016 or TA195, Aug 2010. | In line with NICE TA375 or TA195 criteria or NCL RA pathway | Complete TBF/Blueteq (using agreed local systems) | NHS England is responsible commissioner for paediatric use until transferred to adult services. |
| Infliximab or licensed biosimilar | 10.1 | Cytokine modulators | Psoriatic arthritis | NICE TA199, Aug 2010 | In line with NICE TA199 criteria. | Complete TBF/Blueteq (using agreed local systems) | NHS England is responsible commissioner for paediatric use. |
| Infliximab or licensed biosimilar | 13.3.1 | Cytokine modulators | Psoriasis (adults) | NICE TA134, Jan 2008 | In line with NICE TA134 criteria or NCL Psoriasis pathway (awaited as per JFC) | Complete TBF/Blueteq (using agreed local systems) | NHS England is responsible commissioner for paediatric use. |

| Drug name | BNF section | Category | Licensed Indication/ Indication | NICE Guidance | Indications agreed for funding across NCL | Is this commissioned? What is required for funding approval? | Notes |
|--------------------------------------|-------------------------|---|---|-------------------------------|---|--|---|
| Infliximab or licensed biosimilar | 1.1.3 | Cytokine modulators | Ulcerative colitis (acute exacerbations) | NICE TA163, Dec 2008 | In line with NICE TA163 criteria. | Complete TBF/Blueteq (using agreed local systems) | NHS England is responsible commissioner for paediatric use. |
| Infliximab or licensed biosimilar | 10.1 | Cytokine modulators | Ankylosing spondylitis and non-radiographic axial spondyloarthritis | NICE TA383, Feb 2016. | In line with NICE TA383 criteria. | Complete TBF/Blueteq (using agreed local systems) | NHS England is responsible commissioner for paediatric use. |
| Infliximab or licensed biosimilar | 1.1.3 | Cytokine modulators | Moderately to severely active ulcerative colitis after the failure of conventional therapy | NICE TA329, Feb 2015 | In line with NICE TA329 criteria. | Complete TBF/Blueteq (using agreed local systems) | NHS England is responsible commissioner for paediatric use. |
| Infliximab or licensed biosimilar | 1.1.3 10.1 13.3.1 | Cytokine modulators | All other indications | | NOT COMMISSIONED. | NOT COMMISSIONED. | NHS England is responsible commissioner for paediatric use. NHS England is also commissioner for uveitis, connective tissue disease- interstitial lung disease, graft versus host disease, renal, sarcoidosis and hidradentis suppurativa via the IFR process. NHS England are commissioner for behoets syndrome as per highly specialised criteria only. |
| lxekizumab | 13.3.1 | Humanized anti–interleukin-17 monoclonal antibody | Active psoriatic arthritis in adult patients who have responded inadequately to, or who are intolerant to one or more DMARD therapies | NICE TA537, Aug 2018 | In line with NICE TA 537 criteria. | Complete TBF/Blueteq (using agreed local systems) | |
| lxekizumab | 13.3.1 | Humanized anti–interleukin-17 monoclonal antibody | Moderate to severe plaque psoriasis | NICE TA442, April 2017 | In line with NICE TA442 criteria. | Complete TBF/Blueteq (using agreed local systems) | |
| lxekizumab | 10.1 | Humanized anti–interleukin-17 monoclonal antibody | Ankylosing spondylitis | No NICE guidance anticipated. | NOT COMMISSIONED | NOT COMMISSIONED | |
| lxekizumab | 10.1 | Humanized anti–interleukin-17 monoclonal antibody | Psoriatic arthritis | No NICE guidance anticipated. | NOT COMMISSIONED | NOT COMMISSIONED | |
| Lanreotide | 8.4 | Somatostatin Analogues | All indications except acromegaly, cancer and congenital hyperinsulinism | No NICE guidance anticipated. | Only in line with Local Policies | Complete proforma (using agreed local systems) | NHS England is responsible commissioner for acromegaly, cancer and congenital hyperinsulinism funded by NHS England. |
| Levofloxacin Solution for Inhalation | 5.2 | Antibacterial drugs | Chronic pulmonary infections due to Pseudomonas aeruginosa in adults with CF | No NICE guidance anticipated. | NOT ROUTINELY COMMISSIONED | NOT ROUTINELY COMMISSIONED | NHS England fund through IFR Approval process. |
| Lomitapide | 2.6 | Lipid-regulating drugs | Hypercholesterolaemia, homozygous familial (HoFH). | No NICE guidance anticipated. | NOT ROUTINELY COMMISSIONED | NOT ROUTINELY COMMISSIONED | NHS England fund through IFR Approval process. |
| Mannitol | 3.3.1 | Mucolytics | Indications other than cystic fibrosis | No NICE guidance anticipated. | NOT COMMISSIONED | NOT COMMISSIONED | |
| Mannitol | 3.3.1 | Mucolytics | Cystic Fibrosis | NICE TA266, Nov 2012 | In line with NICE TA266 criteria and NHSE A01/P/b | | NHS England is responsible commissioner. No GP prescribing for paediatric use. |
| Ocriplasmin | 11.6.4 | Retinal disorders | Vitreomacular traction including those associated with macular holes. | NICE TA297, Oct 2013 | In line with NICE TA297 criteria. | Complete proforma (using agreed local systems) | |
| Octreotide | 8.4 | Somatostatin Analogues | All indications except acromegaly, cancer and congenital hyperinsulinism | No NICE guidance anticipated. | Only in line with Local Policies | Complete proforma (using agreed local systems) | NHS England is commissioner for acromegaly, cancer and congenital hyperinsulinism funded by NHS England. |
| Odanacatib | 6.4 | Selective cathepsin-K inhibitor | Postmenopausal osteoporosis | No NICE guidance anticipated. | NOT COMMISSIONED | NOT COMMISSIONED | Development of this drug has been discontinued. |
| Odanacatib | 6.4 | Selective cathepsin-K inhibitor | Osteoporosis in men | No NICE guidance anticipated. | NOT COMMISSIONED | NOT COMMISSIONED | Development of this drug has been discontinued. |
| Omalizumab | 3.1 | Allergen Immunotherapy | Previously treated chronic spontaneous urticaria | NICE TA339, June 2015 | In line with NICE TA339 criteria | Complete TBF/Blueteq (using agreed local systems) | NHS England is responsible commissioner for Asthma. |
| Parenteral Nutrition | 9.4 | Parenteral nutrition- adults | Intestinal failure | NICE CG32. Feb 2006 | Only in line with Local Policies (using agreed systems) | | NHSE is responsible for: Children and Adult specialist intestinal failure services for patients with Intestinal Failure Type II and Type III (including the provision of home parenteral nutrition). |
| Pasireotide | 8.4 | Somatostatin Analogues | Acromegaly | No NICE guidance anticipated. | | NHS England | NHS England is responsible commissioner through IFR approval. |
| Pasireotide | 8.4 | Somatostatin Analogues | Cushings Syndrome (3rd line adults) | No NICE guidance anticipated. | | NHS England | Adults only. GP prescribing not expected |
| Pegaptanib | 11.6.1 | Subfoveal choroidal neovascularisation | Neovascular (wet) age-related macular degeneration | NICE TA155, Aug 2008 | COMMISSIONED | NOT COMMISSIONED | NICE TA155 1.3 Pegaptanib is not recommended for the treatment of wet age-related macular degeneration. |
| Pegpleranib | 11.6.1 | Retinal disorders/intraocular lens replacement surgery | Wet ARMD (age-related macular degeneration) | No NICE guidance anticipated. | NOT COMMISSIONED | NOT COMMISSIONED | Not yet licensed in UK. UK development has been discontinued. |

| Drug name | BNF section | Category | Licensed Indication/ Indication | NICE Guidance | Indications agreed for funding across NCL | Is this commissioned? What is required for funding approval? | Notes |
|----------------------------------|----------------|--|---|----------------------------------|--|--|--|
| Phenylephrine with Ketorolac | 11.4.1 | Retinal disorders / intraocular lens replacement surgery | Intraocular lense replacement surgery | No NICE guidance anticipated. | NOT COMMISSIONED | NOT COMMISSIONED | |
| Ranibizumab | 11.6.1 | Neovascularisation | Neovascular (wet) age-related macular degeneration | NICE TA155, Aug 2008 | In line with NICE crteria TA155 criteria or NCL/NEL wAMD pathway | Complete TBF/Blueteq (using agreed local systems) | |
| Ranibizumab | 11.6.1 | Neovascularisation | Diabetic macular oedema | NICE TA274, Feb 2013 | In line with NICE TA274 criteria or NCL/NEL wAMD pathway | Complete TBF/Blueteq (using agreed local systems) | |
| Ranibizumab | 11.6.1 | Neovascularisation | Macular oedema (retinal vein occlusion) | NICE TA283, May 2013 | | Complete TBF/Blueteq (using agreed local systems) | |
| Ranibizumab | 11.6.1 | Neovascularisation | CNV due to pathological myopia | NICE TA298, Nov 2013 | In line with NICE TA298 criteria or NCL/NEL wAMD pathway | Complete TBF/Blueteq (using agreed local systems) | |
| Riluzole | 10.3 | Torsion Dystonia and other involuntary movements | ALS form of Motor Neurone Disease | NICE TA20, Jan 2001 | In line with NICE TA20 criteria | Complete TBF/Blueteq (using agreed local systems) | Adults only. |
| Rituximab or licensed biosimilar | 10.1 | Cytokine modulators | Rheumatoid arthritis after failure of TNF inhibitor | NICE TA195, Aug 2010 | In line with NICE TA195 criteria | Complete TBF/Blueteq (using agreed local systems) | Adults only, GP prescribing not expected. NHS England is responsible commissioner for cancerous and specialist auto-inflammatotory conditions. |
| Rituximab or licensed biosimilar | 10.1 | Cytokine modulators | Rheumatoid arthritis (first line) | No NICE guidance anticipated. | See JFC pathway | NOT COMMISSIONED | Adults only. NHS England is responsible commissioner for cancerous and specialist auto-inflammatotory conditions. |
| Rituximab or licensed biosimilar | N/A | Cytokine modulators | Autoimmune Haemolytic Anaemia (AIHA). | No NICE guidance anticipated. | In line with JFC: For AIHA. July 2016 | Notification through SLAM (using agreed local systems) | Adults only. NHS England is responsible commissioner for cancerous and specialist auto-inflammatotory conditions. |
| Rituximab or licensed biosimilar | N/A | Cytokine modulators | ÎTP | No NICE guidance anticipated. | In line with JFC: For refractory or relapsed disease only (November 2012) | Complete TBF/Blueteq (using agreed local systems) | Adults only. NHS England is responsible commissioner for cancerous and specialist auto-inflammatotory conditions. |
| Rituximab or licensed biosimilar | 10.1 | Cytokine modulators | ANCA-positive vasculitis | NICE TA308, Mar 2014 | In line with NICE TA 308 and NHS England policy A13/P/a | NHS England | Adults only.No GP prescribing. NHS England is responsible commissioner for cancerous and specialist auto- inflammatotory conditions. |
| Rituximab or licensed biosimilar | 10.1 | Cytokine modulators | Juvenile Arthritis | NHS England Policy | In line with NHS England policy: E03/P/d | NHS England | No GP prescribing. NHS England is responsible commissioner for cancerous and specialist auto-inflammatotory conditions. |
| Rituximab or licensed biosimilar | N/A | Cytokine modulators | Neuromyelitis optica | NHS England | In line with NHS England specification | NHS England | The first 2 courses of Rituximab, i.e. 1st month and 6th month (1 course-2 injections on day 1 and day 14) will be given and funded by NHS England commissioned centres. The central budget for Rituximab is held by The Walton Centre NHS Foundation Trust and that Oxford will cross charge Rituximab to the Walton Centre based on use. The centres will assess benefit after the second course of treatment and if ongoing care is needed Rituximab may be given locally as part of a shared care arrangement. On-going drug funding will be the responsibility of the patient's CCCG. |
| Rituximab or licensed biosimilar | 10.1 | Cytokine modulators | Other indications (adults) used in CCG commissioned service | | | NHS England responsibility | Adults only.No GP prescribing. NHS England is responsible commissioner for cancerous and specialist auto- linflammatotory conditions. |
| Romiplostim | 9.4.2 | Platelet Disorder Drugs | Chronic immune (idiopathic) thrombocytopenic purpura | NICE TA221, April 2011 | In line with NICE TA221 criteria | Complete TBF/Blueteq (using agreed local systems) | Adults only. GP prescribing NOT expected. NHS England is responsible commissioner for paediatric use. |
| Sarilumab | 10.1 | Cytokine modulators | Rheumatoid arthritis | NICE TA485, Nov 2017 | In line with NICE TA485 criteria. | Complete TBF/blueteq (using agreed local systems) | Adults only. GP prescribing NOT expected. NHS England responsible commissioner for paediatric use through IFR process. |
| Secukinumab | 10.1 | Cytokine modulators | Ankylosing spondylitis after treatment with non-steroidal anti-inflammatory drugs or TNF-alpha inhibitors | NICE TA407, Sept 2016 | In line with NICE TA407 criteria. | Complete TBF/Blueteq (using agreed local systems) | Adults only. GP prescribing NOT expected. NHS England responsible commissioner for paediatric use through IFR process. |
| Secukinumab | 13.3.1 | Cytokine modulators | Moderate to severe plaque psoriasis | NICE TA350, July 2015. | TA350 criteria. | Complete TBF/Blueteq (using agreed local systems) | Adults only. GP prescribing NOT expected. NHS England responsible commissioner for paediatric use through IFR process. |
| Secukinumab | 10.1 | Cytokine modulators | Active psoriatic arthritis after DMARDs | NICE TA445, May 2017 | In line with NICE TA445 criteria. | Complete TBF/Blueteq (using agreed local systems) | Adults only. GP prescribing NOT expected. NHS England responsible commissioner for paediatric use through IFR process. |
| Secukinumab | 10.1 | Cytokine modulators | Rheumatoid arthritis | No NICE guidance anticipated. | NOT COMMISSIONED | NOT COMMISSIONED | Not yet licensed in UK. NHS England responsible for paediatric indication |
| Serelaxin | N/A | Hypertension and Heart failure | Heart failure | NICE Guidance | NOT COMMISSIONED | NOT COMMISSIONED | Not yet licensed in UK. UK development has been discontinued. Appraisal suspended indefinitely. |
| Siltuximab | 10.1 | Cytokine modulators | Castleman's disease | No NICE guidance anticipated. | | NOT COMMISSIONED | |

| Drug name | BNF section | Category | Licensed Indication/ Indication | NICE Guidance | Indications agreed for funding across NCL | Is this commissioned? What is required for funding approval? | Notes |
|---|----------------|--|--|---|--|--|--|
| Sirukumab | 10.1 | Cytokine modulators | Rheumatoid arthritis | No NICE guidance anticipated. | | NOT COMMISSIONED | Company decided to withdraw its marketing authorisation application. Consequently, appraisal suspended in October 2017. |
| Sodium oxybate | 4.7.2 | Hypnotics and anxiolytics | Narcolepsy with cataplexy (adults) | No NICE guidance anticipated. | local policy being updated | Complete using agreed local systems | NHS England is responsible commissioner for paediatric services in line with clinical commissioning policy 16065/P. |
| Somatropin (adults) | 6.7.4 | Growth Hormone or Growth Hormone Receptor Antagonist | Growth hormone deficiency | NICE TA64, Aug 2003 | In line with NICE TA64 criteria. | Complete TBF/Blueteq (using agreed local systems) | NHS England is responsible for indications falling outside NICE guidance treated in specialist centres |
| Somatropin (children) | 6.7.4 | Growth Hormone or Growth Hormone Receptor Antagonist | Growth failure in children | NICE TA188, May 2010 | In line with NICE TA188 criteria. | Complete TBF/Blueteq (using agreed local systems) | NHS England is responsible for indications falling outside NICE guidance treated in specialist centres |
| Tafamidis | 4.4.1 | Neuroprotective agents | Treatment of transthyretin amyloidosis in adult patients with stage 1 symptomatic polyneuropathy to delay peripheral neurologic impairment | N/A | NOT COMMISSIONED | NOT COMMISSIONED | Current development status of this drug is unknown |
| Teriflunomide | 8.1.1 | Other immunomodulating drugs | Relapsing-remitting multiple sclerosis | NICE TA303, Jan 2014 and NHS England Policy | In line with NICE TA303 criteria and NHS England Policy: D04/P/b | NHS England | NHS England is responsible commissioner. |
| Teriparatide | 6.4 | Drugs affecting bone metabolism | Secondary prevention of osteoporotic fragility fractures in postmenopausal women | NICE TA161, Oct 2008 | In line with NICE TA161 criteria | Complete TBF/Blueteq (using agreed local systems) | |
| Teriparatide | 6.4 | Drugs affecting bone metabolism | Treatment of osteoporosis associated with glucocorticoids | No NICE guidance anticipated. | NOT COMMISSIONED | NOT COMMISSIONED | |
| Teriparatide | 6.4 | Drugs affecting bone metabolism | Osteogenesis imperfecta | NHS England Policy | Not routinely commissioned. In line with NICE policy 16002/P | NHS England | NHS England are the responsible commissioner through the IFR process. |
| Teriparatide | 6.4 | Drugs affecting bone metabolism | Male and juvenile osteoporosis | NHS England | Not routinely commissioned. In line with IFR approval. | NHS England | NHS England are the responsible commissioner through the IFR process. |
| Tesamorelin | N/A | Growth Hormone & growth hormone Receptor Antagonist | HIV associated lipodystrophy | No NICE guidance anticipated. | NOT COMMISSIONED | NOT COMMISSIONED | Filing for this drug has been withdrawn |
| Thrombomodulin, Recombinant Human | N/A | Fibrinolytics | Septic shock (Sepsis) (adult, non-specialist services) | No NICE guidance anticipated. | NOT COMMISSIONED | NOT COMMISSIONED | Launch expected 2019 |
| Tildrakizumab | N/A | Skin conditions | Psoriasis (adults) | No NICE guidance. ID1060 | NOT COMMISSIONED | NOT COMMISSIONED | Expected launch April 2019 |
| Tobramycin (Tobi® / Bramitob®) nebuliser solution | 5.2 | Antibacterial Drugs | Cystic fibrosis | | In line with NICE TA276 criteria & NHS England Policy:A01/P/b | NHS England | NHS England is the responsible commissioner. |
| Tobramycin (Tobi® Podhaler) dry powder for inhalation | 5.2 | Antibacterial Drugs | Cystic fibrosis | NICE TA276, Mar 2013. | In line with NICE TA276 criteria & NHS England Policy:A01/P/b | NHS England | NHS England is the responsible commissioner. |
| Tocilizumab | 10.1 | Cytokine modulator | Rheumatoid arthritis | NICE TA247, Feb 2012 or TA375, Jan 2016 | In line with NICE crteria TA247 or TA375 criteria. | Complete TBF/Blueteq (using agreed local systems) | Adults only, GP prescribing NOT expected. NHS England is responsible commissioner for paediatric use until transferred to adult service. |
| Tocilizumab | 10.1 | Cytokine modulator | Juvenile arthritis | NICE TA373, Dec 2015 and NHS England commissioning Policy | In line with NICE TA373 and NHS England commissioning policy E03/P/d | NHS England | NHS England is responsible commissioner for juvenile arthritis in paediatrics, until transferred into adult service. |
| Tocilizumab | 10.1 | Cytokine modulator | Other indications (adults) used in CCG commissioned service | No NICE guidance anticipated. | NOT COMMISSIONED | NOT COMMISSIONED | |
| Tocilizumab | 10.1 | Cytokine modulator | Takayasu arteritis | NHS England clinical commissioning | In line with NHS England commissioning policy:16056/P | NHS England | NHS England is the responsible commissioner. |
| Tocilizumab | 10.1 | Cytokine modulator | Giant Cell Arteritis | NICE TA518, Apr 2018 and NHS England Policy. | Not routinely commissioned. In line | Not routinely commissioned. NHS England. | NHS England is the responsible commissioner through the IFR process. |
| Tofacitinib | 10.1 | Cytokine modulator | Moderate to severe rheumatoid arthritis | NICE TA480, Oct 2017 | In line with NICE TA480 criteria. | Complete TBF/Blueteq (using agreed local systems) | Adults only, GP prescribing NOT expected. NHS England is responsible commissioner for paediatric use through the IFR process. |

(Collaboration Commissioners' list of Indications that may be commissioned for excluded drugs 2019-20)

| Drug name | BNF section | Category | Licensed Indication/ Indication | NICE Guidance | NCL | Is this commissioned? What is required for funding approval? | Notes |
|-------------|----------------|--|--|----------------------------------|--|--|---|
| Tofacitinib | 10.1 | Cytokine modulator | Treating active psoriatic arthritis after inadequate response to DMARDs | NICE TA543, Oct 2018 | In line with NICE TA543 criteria. | Complete TBF/Blueteq (using agreed local systems) | Adults only, GP prescribing NOT expected. NHS England is responsible commissioner for paediatric use through the IFR process. |
| Tofacitinib | 1.1.3 | Cytokine modulator | Moderately to severely active ulcerative colitis | NICE TA547, Nov 2018 | In line with NICE TA547 criteria. | Complete TBF/Blueteq (using agreed local systems) | Adults only. GP prescribing NOT expected. NHS England is responsible commissioner for paediatric use through the IFR process. |
| Tofacitinib | 13.3.1 | Cytokine modulator | Psoriasis (adults) | No NICE guidance anticipated. | NOT COMMISSIONED | NOT COMMISSIONED | UK development for this has been discontinued. |
| Tofacitinib | 10.1 | Cytokine modulator | Juvenile arthritis | No NICE guidance anticipated. | NOT COMMISSIONED | NOT COMMISSIONED | Adults only. GP prescribing NOT expected. NHS England is responsible commissioner for paediatric use through the IFR process. |
| Tolvaptan | 6.1.2 | Vasopressin V2- receptor antagonist | Autosomal dominant polycystic kidney disease, stage 2 or 3, rapidly progressing | NICE TA358, Oct 2015 | In line with NICE TA358 criteria. | Complete TBF/Blueteq (using agreed local systems) | No GP prescribing. NHS England is responsible commissioner for hyponatraemia in cancer. |
| Tolvaptan | 6.1.2 | Vasopressin V2- receptor antagonist | Hyponatraemia due to syndrome of inappropriate antidiuretic hormone secretion (SIADH). | No NICE guidance anticipated. | NOT COMMISSIONED | NOT COMMISSIONED | No GP prescribing. NHS England is responsible commissioner for hyponatraemia in cancer. |
| Tolvaptan | 6.1.2 | Vasopressin V2- receptor antagonist | Hyponatraemia from other causes and other endocrine uses | No NICE guidance anticipated. | NOT COMMISSIONED | NOT COMMISSIONED | No GP prescribing. NHS England is responsible commissioner for hyponatraemia in cancer. |
| Ustekinumab | 13.3.1 | Drugs affecting the immune response | Psoriasis moderate to severe | NICE TA180, Sept 2009. | In line with NICE crteria TA180 criteria or NCL Psoriasis pathway (awaited as per JFC) | Complete TBF/Blueteq (using agreed local systems) | Adults only. GP prescribing NOT expected. NHS England is responsible commissioner for paediatric use. |
| Ustekinumab | 1.1.3 | Drugs affecting the immune response | Crohn's disease | NICE TA456, July 2017 | In line with NICE TA456 criteria. | Complete TBF/Blueteq (using agreed local systems) | Adults only. GP prescribing NOT expected. NHS England is responsible commissioner for paediatric use. |
| Ustekinumab | 10.1 | Drugs affecting the immune response | Active psoriatic arthritis | NICE TA340, June 2015 | In line with NICE TA340 criteria. | Complete TBF/Blueteq (using agreed local systems) | Adults only. GP prescribing NOT expected. NHS England is responsible commissioner for paediatric use. |
| Ustekinumab | 13.3.1 | Drugs affecting the immune response | Plaque psoriasis in children and young people | NICE TA455, July 2017 | In line with NICE TA455 criteria. | NHS England | NHS England is responsible commissioner. |
| Vedolizumab | 1.1.3 | Drugs affecting the immune response | Crohn's disease second line after conventional therapy and TNF inhibitors | | In line with NICE | Complete TBF/Blueteq (using agreed local systems) | Adults only. GP prescribing NOT expected. NHS England is responsible commissioner for paediatric use. |
| Vedolizumab | 1.1.3 | Drugs affecting the immune response | Moderately to severely active ulcerative colitis | NICE TA342, June 2015 | In line with NICE TA342 criteria. | Complete TBF/Blueteq (using agreed local systems) | Adults only. GP prescribing NOT expected. NHS England is responsible commissioner for paediatric use. |
| Verteporfin | 11.6.1 | Subfoveal choroidal neovascularisation | Photodynamic therapy for wet age-related macular degeneration. | NICE TA68, Sept 2003 | In line with local policies | Complete TBF/Blueteq (using agreed local systems) | No GP prescribing |

Last reviewed: March 2019