

Formulary Adherence Checklist for NICE Technology Appraisals About Medicines



Developed for the NHS by: **East and South East England Specialist Pharmacy Services**

Milton Keynes Health Economy

This spreadsheet is updated monthly and enables self-audit of a medicines formulary for adherence to current NICE Technology Appraisals. **Version 20 November 2017**

All guidelines refer to adults unless indicated. No copyright is asserted on this material if used for non-commercial purposes within the NHS.

Technology appraisal (TA) Titles are hyperlinks to full guidance	Date of TA Release	Availability of medicine for NHS patients with this medical condition, as indicated by NICE	Adherence of local formulary to NICE				
			Yes (mark 'x' if applicable)	N/A (mark 'x' if applicable)	Date of local decision (DD/MM/YY)	Time to implement (days)	Notes (e.g. rationale, method of making available)
2017/2018							
Venetoclax for treating chronic lymphocytic leukaemia (TA487)	01/10/2017	Venetoclax (Venclyxto®) is recommended for use within the Cancer Drugs Fund, within its marketing authorisation, as an option for treating chronic lymphocytic leukaemia	x		29/11/2017	59	Approved for addition to the formulary and to be used in line with NICE guidance and commissioning statements. Cancer Drug Fund
Aflibercept for treating choroidal neovascularisation (TA486)	01/10/2017	Aflibercept (Eylea®) is recommended, within its marketing authorisation, as an option for treating visual impairment because of myopic choroidal neovascularisation in adults	x		29/11/2017	59	Commissioning responsibility: CCG; Approved for addition to the formulary and to be used in line with NICE guidance and commissioning statements
Sarilumab for moderate to severe rheumatoid arthritis (TA485)	01/10/2017	Sarilumab (Kevzara®) with methotrexate, is recommended as an option for treating active rheumatoid arthritis in adults whose disease has responded inadequately to intensive therapy with a combination of conventional disease-modifying anti-rheumatic drugs (DMARDs)	x		29/11/2017	59	Commissioning responsibility: CCG; Approved for addition to the formulary and to be used in line with NICE guidance and commissioning statements
Nivolumab for previously treated non-squamous non-small-cell lung cancer (TA484)	01/10/2017	Nivolumab (Opdivo®) is recommended for use within the Cancer Drugs Fund as an option for treating locally advanced or metastatic non-squamous non-small-cell lung cancer in adults after chemotherapy	x		29/11/2017	59	Approved for addition to the formulary and to be used in line with NICE guidance and commissioning statements. Cancer Drug Fund
Nivolumab for previously treated squamous non-small-cell lung cancer (TA483)	01/10/2017	Nivolumab (Opdivo®) is recommended for use within the Cancer Drugs Fund as an option for treating locally advanced or metastatic squamous non-small-cell lung cancer in adults after chemotherapy	x		29/11/2017	59	Approved for addition to the formulary and to be used in line with NICE guidance and commissioning statements. Cancer Drug Fund
Immunosuppressive therapy for kidney transplant in children and young people (TA482)	01/10/2017	Immunosuppressive therapy for kidney transplant in children and young people	x		29/11/2017	59	Remove NICE technology appraisal guidance 85. Add the immunosuppressive drugs to the formulary and link NICE TA to each drug; To add both the positive and negative statements to the formulary
Immunosuppressive therapy for kidney transplant in adults (TA481)	01/10/2017	Immunosuppressive therapy for kidney transplant in adults	x		29/11/2017	59	Remove NICE technology appraisal guidance 85. Add the immunosuppressive drugs to the formulary and link NICE TA to each drug; To add both the positive and negative statements to the formulary

Tofacitinib for moderate to severe rheumatoid arthritis (TA480)	01/10/2017	Tofacitinib (Xeljanz®) is recommended as an option for treating active rheumatoid arthritis in adults whose disease has responded inadequately to intensive therapy with a combination of conventional disease-modifying anti-rheumatic drugs (DMARDs)	x		29/11/2017	59	Commissioning responsibility: CCG; PBR Excluded. Discontinued Patient Access, Hospital Prescribing Only. Approved for addition to the formulary and to be used in line with NICE guidance and commissioning statements
Reslizumab for treating severe eosinophilic asthma (TA479)	01/10/2017	Reslizumab (Cinqaero®) is recommended as an option for the treatment of severe eosinophilic asthma that is inadequately controlled in adults despite maintenance therapy with high-dose inhaled corticosteroids plus another drug	x		29/11/2017	59	Approved for addition to the formulary and to be used in line with NICE guidance and commissioning statements
Brentuximab vedotin for treating relapsed or refractory systemic anaplastic large cell lymphoma (TA478)	01/10/2017	Brentuximab vedotin (Adcetris®) is recommended as an option for treating relapsed or refractory systemic anaplastic large cell lymphoma in adults	x		29/11/2017	59	Approved for addition to the formulary and to be used in line with NICE guidance and commissioning statements
Autologous chondrocyte implantation for treating symptomatic articular cartilage defects of the knee (TA477)	01/10/2017	Autologous chondrocyte implantation (ACI) ChondroCelect® is recommended as an option for treating symptomatic articular cartilage defects of the knee	x		29/11/2017	59	NICE Appraisal N/A at MK
Paclitaxel as albumin-bound nanoparticles with gemcitabine for untreated metastatic pancreatic cancer (TA476)	06/09/2017	Paclitaxel as albumin-bound nanoparticles (nab-paclitaxel) with gemcitabine is recommended as an option for untreated metastatic adenocarcinoma of the pancreas in adults	x		27/09/2017	21	Approved for addition to the formulary and to be used in line with NICE guidance and commissioning statements
Dimethyl fumarate for treating moderate to severe plaque psoriasis (TA475)	06/09/2017	Dimethyl fumarate is recommended as an option for treating plaque psoriasis in adults	x		29/11/2017	84	Confirmation of place in therapy is required. Agreed to add in line with Oxford Protocol. Issues to be resolved outside of meeting by 6th December 2017
Sorafenib for treating advanced hepatocellular carcinoma (TA474)	06/09/2017	Sorafenib is recommended as an option for treating advanced hepatocellular carcinoma only for people with Child-Pugh grade A liver impairment	x		27/09/2017	21	Approved for addition to the formulary and to be used in line with NICE guidance and commissioning statements
Cetuximab for treating recurrent or metastatic squamous cell cancer of the head and neck (TA473)	31/08/2017	Cetuximab in combination with platinum-based chemotherapy is recommended as an option for treating recurrent or metastatic squamous cell cancer of the head and neck in adults	x		27/09/2017	27	CCG commissioned. Approved for addition to the formulary and to be used in line with NICE guidance and commissioning statements.
Obinutuzumab with bendamustine for treating follicular lymphoma refractory to rituximab (TA472)	30/08/2017	Obinutuzumab in combination with bendamustine followed by obinutuzumab maintenance is recommended for use within the Cancer Drugs Fund as an option for treating adults with follicular lymphoma that did not respond or progressed during or up to 6 months after treatment with rituximab or a rituximab-containing regimen	x		27/09/2017	28	Approved for addition to the formulary and to be used in line with NICE guidance and commissioning statements
Eluxadoline for treating irritable bowel syndrome with diarrhoea (TA471)	30/08/2017	Eluxadoline is recommended as an option for treating irritable bowel syndrome with diarrhoea in adults	x		27/09/2017	28	Approved for addition to the formulary and to be used in line with NICE guidance with Secondary Care Initiation and stabilisation including one month review.

Ofatumumab with chemotherapy for treating chronic lymphocytic leukaemia (terminated appraisal) (TA470)	23/08/2017	Ofatumumab with chemotherapy - was unable to make a recommendation about the use in the NHS of ofatumumab with chemotherapy for treating chronic lymphocytic leukaemia because no evidence submission was received from Novartis Pharmaceuticals UK	x		27/09/2017	35	Terminated Appraisal by NICE
Idelalisib with ofatumumab for treating chronic lymphocytic leukaemia (terminated appraisal) (TA469)	23/08/2017	NICE is unable to make a recommendation about the use in the NHS of idelalisib with ofatumumab for treating chronic lymphocytic leukaemia because no evidence submission was received	x		27/09/2017	35	Terminated Appraisal by NICE
Methylnaltrexone bromide for treating opioid-induced constipation (terminated appraisal) (TA468)	23/08/2017	Methylnaltrexone bromide - was unable to make a recommendation about the use in the NHS of methylnaltrexone bromide for treating opioid-induced constipation because no evidence submission was received from Swedish Orphan Biovitrum Ltd	x		27/09/2017	35	Terminated Appraisal by NICE
Holoclar for treating limbal stem cell deficiency after eye burns (TA467)	16/08/2017	Holoclar (ex vivo expanded autologous human corneal epithelial cells containing stem cells) is recommended as an option in people with moderate to severe limbal stem cell deficiency after eye burns	x		27/09/2017	42	Approved for addition to the formulary and to be used in line with NICE guidance and commissioning statements
Baricitinib for moderate to severe rheumatoid arthritis (TA466)	09/08/2017	Baricitinib, with methotrexate is recommended as an option for treating active rheumatoid arthritis in adults whose disease has responded inadequately to intensive therapy with a combination of conventional disease-modifying antirheumatic drugs (DMARDs),	x		27/09/2017	49	Approved for addition to the formulary and to be used in line with NICE guidance and commissioning statements
Olaratumab in combination with doxorubicin for treating advanced soft tissue sarcoma (TA465)	09/08/2017	Olaratumab, in combination with doxorubicin , is recommended for use within the Cancer Drugs Fund as an option for advanced soft tissue sarcoma in adults	x		27/09/2017	49	Approved for addition to the formulary and to be used in line with NICE guidance and commissioning statements
Bisphosphonates for treating osteoporosis (TA464)	09/08/2017	Oral bisphosphonates (alendronic acid, ibandronic acid and risedronate sodium) are recommended as options for treating osteoporosis in adults	x		27/09/2017	49	Approved for addition to the formulary and to be used in line with NICE guidance and commissioning statements
Cabozantinib for previously treated advanced renal cell carcinoma (TA463)	09/08/2017	Cabozantinib is recommended, within its marketing authorisation, as an option for treating advanced renal cell carcinoma in adults after vascular endothelial growth factor (VEGF)-targeted therapy	x		27/09/2017	49	Approved for addition to the formulary and to be used in line with NICE guidance and commissioning statements
Nivolumab for treating relapsed or refractory classical Hodgkin lymphoma (TA462)	26/07/2017	Nivolumab is recommended, within its marketing authorisation, as an option for treating relapsed or refractory classical Hodgkin lymphoma in adults after autologous stem cell transplant and treatment with brentuximab vedotin, when the company provides nivolumab with the discount agreed in the patient access scheme.	x		27/09/2017	63	Approved for addition to the formulary and to be used in line with NICE guidance and commissioning statements

Roflumilast for treating chronic obstructive pulmonary disease (TA461)	26/07/2017	Roflumilast , as an add-on to bronchodilator therapy, is recommended as an option for treating severe chronic obstructive pulmonary disease in adults with chronic bronchitis	x		27/09/2017	63	Approved for addition to the formulary and to be used in line with NICE guidance and commissioning statements
Adalimumab and dexamethasone for treating non-infectious uveitis (TA460)	26/07/2017	Adalimumab is recommended as an option for treating non-infectious uveitis in the posterior segment of the eye in adults with inadequate response to corticosteroids. Dexamethasone (Ozurdex®) intravitreal implant is recommended as an option for treating non-infectious uveitis in the posterior segment of the eye in adults	x		27/09/2017	63	Approved for addition to the formulary and to be used in line with NICE guidance and commissioning statements
Collagenase clostridium histolyticum for treating Dupuytren's contracture (TA459)	26/07/2017	Collagenase clostridium histolyticum for people who meet the inclusion criteria for the ongoing clinical trial (HTA-15/102/04), comparing collagenase clostridium histolyticum (CCH) with limited fasciectomy, are encouraged to participate in the study. For people not taking part in the ongoing clinical trial, CCH is recommended as an option for treating Dupuytren's contracture with a palpable cord in adults	x		27/09/2017	63	Approved for addition to the formulary and to be used in line with NICE guidance and commissioning statements
Trastuzumab emtansine for treating HER2-positive advanced breast cancer after trastuzumab and a taxane (TA458)	19/07/2017	Trastuzumab emtansine is recommended, within its marketing authorisation, as an option for treating human epidermal growth factor receptor 2 (HER2)-positive, unresectable, locally advanced or metastatic breast cancer in adults who previously received trastuzumab and a taxane, separately or in combination.	x		27/09/2017	70	Approved for addition to the formulary and to be used in line with NICE guidance and commissioning statements
Carfilzomib for previously treated multiple myeloma (TA457)	19/07/2017	Carfilzomib in combination with dexamethasone is recommended as an option for treating multiple myeloma in adults	x		27/09/2017	70	Approved for addition to the formulary and to be used in line with NICE guidance and commissioning statements
Ustekinumab for moderately to severely active Crohn's disease after previous treatment (TA456)	12/07/2017	Ustekinumab is recommended, within its marketing authorisation, as an option for treating moderately to severely active Crohn's disease, that is, for adults who have had an inadequate response with, lost response to, or were intolerant to either conventional therapy or a TNF-alpha inhibitor or have medical contraindications to such therapies.	x		27/09/2017	77	Approved for addition to the formulary and to be used in line with NICE guidance and commissioning statements
Adalimumab, etanercept and ustekinumab for treating plaque psoriasis in children and young people (TA455)	12/07/2017	Adalimumab, Etanercept and Ustekinumab is recommended as an option for treating plaque psoriasis in children and young people aged 4 years or older	x		27/09/2017	77	Approved for addition to the formulary and to be used in line with NICE guidance and commissioning statements

Daratumumab with lenalidomide and dexamethasone for treating relapsed or refractory multiple myeloma (terminated appraisal) (TA454)	05/07/2017	Daratumumab - was unable to make a recommendation about the use in the NHS of daratumumab, with lenalidomide and dexamethasone, for treating relapsed or refractory multiple myeloma because no evidence submission was received from Janssen-Cilag	x		26/07/2017	21	Terminated Appraisal by NICE
Bortezomib for treating multiple myeloma after second or subsequent relapse (terminated appraisal) (TA453)	05/07/2017	Bortezomib - was unable to make a recommendation about the use in the NHS of bortezomib for treating multiple myeloma after second or subsequent relapse because no evidence submission was received from Janssen-Cilag	x		26/07/2017	21	Terminated Appraisal by NICE
Ibrutinib for untreated chronic lymphocytic leukaemia without a 17p deletion or TP53 mutation (terminated appraisal) (TA452)	05/07/2017	Ibrutinib - was unable to make a recommendation about the use in the NHS of ibrutinib for untreated chronic lymphocytic leukaemia without a 17p deletion or TP53 mutation because no evidence submission was received from Janssen-Cilag	x		26/07/2017	21	Terminated Appraisal by NICE
Ponatinib for treating chronic myeloid leukaemia and acute lymphoblastic leukaemia (TA451)	28/06/2017	Ponatinib (Iclusig) is recommended, within its marketing authorisation, as an option for treating chronic-, accelerated- or blast-phase chronic myeloid leukaemia in adults	x		26/07/2017	28	NHSE commissioned. Approved for addition to the formulary and to be used in line with NICE guidance and commissioning statements
Blinatumomab for previously treated Philadelphia-chromosome-negative acute lymphoblastic leukaemia (TA450)	28/06/2017	Blinatumomab (Blincyto) is recommended within its marketing authorisation as an option for treating Philadelphia-chromosome-negative relapsed or refractory precursor B-cell acute lymphoblastic leukaemia in adults, only if the company provides it with the discount agreed in the patient access scheme.	x		26/07/2017	28	NHSE commissioned. Approved for addition to the formulary and to be used in line with NICE guidance and commissioning statements
Everolimus and sunitinib for treating unresectable or metastatic neuroendocrine tumours in people with progressive disease (TA449)	28/06/2017	Everolimus (Afinitor) and Sunitinib (Sutent) are recommended, within their marketing authorisations, as options for treating well- or moderately differentiated unresectable or metastatic neuroendocrine tumours (NETs) of pancreatic origin in adults with progressive disease.	x		26/07/2017	28	NHSE commissioned. Approved for addition to the formulary and to be used in line with NICE guidance and commissioning statements
Etelcalcetide for treating secondary hyperparathyroidism (TA448)	28/06/2017	Etelcalcetide (Parsabiv) is recommended as an option for treating secondary hyperparathyroidism in adults with chronic kidney disease on haemodialysis	x		26/07/2017	28	NHSE commissioned. Approved for addition to the formulary and to be used in line with NICE guidance and commissioning statements
Pembrolizumab for untreated PD-L1-positive metastatic non-small-cell lung cancer (TA447)	28/06/2017	Pembrolizumab (Keytruda) is recommended for use within the Cancer Drugs Fund as an option for untreated PD-L1-positive metastatic non-small-cell lung cancer in adults	x		26/07/2017	28	NHSE commissioned. Approved for addition to the formulary and to be used in line with NICE guidance and commissioning statements
Brentuximab vedotin for treating CD30-positive Hodgkin lymphoma (TA446)	28/06/2017	Brentuximab vedotin (Adcetris) is recommended as an option for treating CD30-positive Hodgkin lymphoma in adults	x		26/07/2017	28	NHSE commissioned. Approved for addition to the formulary and to be used in line with NICE guidance and commissioning statements

Certolizumab pegol and secukinumab for treating active psoriatic arthritis after inadequate response to DMARDs (TA445)	24/05/2017	Certolizumab pegol (Cimzia) alone, or in combination with methotrexate is recommended as an option for treating active psoriatic arthritis in adults	x		26/07/2017	63	CCG commissioned. Approved for addition to the formulary and to be used in line with NICE guidance and commissioning statements.
Afatinib for treating advanced squamous non-small-cell lung cancer after platinum-based chemotherapy (terminated appraisal) (TA444)	22/03/2017	Tenofovir alafenamide - was unable to make a recommendation about the use in the NHS of afatinib for treating locally advanced or metastatic squamous non-small-cell lung cancer after platinum-based chemotherapy because no evidence submission was received from Boehringer Ingelheim.	x		24/05/2017	63	Terminated Appraisal by NICE
Obeticholic acid for treating primary biliary cholangitis (TA443)	26/04/2017	Obeticholic acid (Ocaliva) is recommended, within its marketing authorisation, as an option for treating primary biliary cholangitis in combination with ursodeoxycholic acid for people whose disease has responded inadequately to ursodeoxycholic acid or as monotherapy for people who cannot tolerate ursodeoxycholic acid. Obeticholic acid is recommended only if the company provides it with the discount agreed in the patient access scheme.	x		24/05/2017	28	NHSE commissioned. Approved for addition to the formulary and to be used in line with NICE guidance and commissioning statements
Ixekizumab for treating moderate to severe plaque psoriasis (TA442)	26/04/2017	Ixekizumab (Taltz) is recommended as an option for treating plaque psoriasis in adults	x		24/05/2017	28	CCG commissioned. Approved for addition to the formulary and to be used in line with NICE guidance and commissioning statements.
Daclizumab for treating relapsing–remitting multiple sclerosis (TA441)	26/04/2017	Daclizumab (Zinbryta) is recommended as an option for treating multiple sclerosis in adults	x		24/05/2017	28	NHSE commissioned. Approved for addition to the formulary and to be used in line with NICE guidance and commissioning statements
Pegylated liposomal irinotecan for treating pancreatic cancer after gemcitabine (TA440)	26/04/2017	Pegylated liposomal irinotecan (Onivyde) , in combination with 5-fluorouracil and leucovorin, is not recommended, within its marketing authorisation, for treating metastatic adenocarcinoma of the pancreas in adults whose disease has progressed after gemcitabine-based therapy.	x		24/05/2017	28	Not recommended
			49	0			
			% "Yes"	% "N/A"	-		Average implement time (days)
Adherence statistics for 2017-18			100%	0%			46