

## 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 March 2018**

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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)
<b>2017/2018</b>							
<a href="#">Pertuzumab with trastuzumab and docetaxel for treating HER2-positive breast cancer (TA509)</a>	07/03/2018	<b>Pertuzumab (Perjeta®) in combination with trastuzumab and docetaxel</b> , is recommended, within its marketing authorisation, for treating HER2-positive metastatic or locally recurrent unresectable breast cancer, in adults who have not had previous anti-HER2 therapy or chemotherapy for their metastatic disease	x		28/03/2018	21	NHSE commissioned. Approved for addition to the formulary and to be used in line with NICE guidance and commissioning statements and the National Cancer Drugs Fund requirements
<a href="#">Autologous chondrocyte implantation using chondrosphere for treating symptomatic articular cartilage defects of the knee (TA508)</a>	07/03/2018	<b>Autologous chondrocyte implantation (ACI) using chondrosphere (Spherox®)</b> is recommended as an option for treating symptomatic articular cartilage defects of the femoral condyle and patella of the knee (International Cartilage Repair Society grade III or IV) in adults	x		28/03/2018	21	NHSE commissioned. Approved for addition to the formulary and to be used in line with NICE guidance and commissioning statements
<a href="#">Sofosbuvir–velpatasvir–voxilaprevir for treating chronic hepatitis C (TA507)</a>	21/02/2018	<b>sofosbuvir–velpatasvir–voxilaprevir (Vosevi®)</b> is recommended as an option for treating chronic hepatitis C in adults	x		28/03/2018	35	NHSE commissioned. Approved for addition to the formulary and to be used in line with NICE guidance and commissioning statements
<a href="#">Lesinurad for treating chronic hyperuricaemia in people with gout (TA506)</a>	07/02/2018	<b>Lesinurad (Zurampic®)</b> is not recommended within its marketing authorisation, that is, with a xanthine oxidase inhibitor for treating hyperuricaemia in adults with gout whose serum uric acid is above the target level despite an adequate dose of a xanthine oxidase inhibitor alone	x		28/03/2018	49	Not recommended
<a href="#">Ixazomib with lenalidomide and dexamethasone for treating relapsed or refractory multiple myeloma (TA505)</a>	07/02/2018	<b>Ixazomib (Ninlaro®) with lenalidomide and dexamethasone</b> is recommended for use within the Cancer Drugs Fund as an option for treating multiple myeloma in adults	x		28/03/2018	49	NHSE commissioned. Approved for addition to the formulary and to be used in line with NICE guidance and commissioning statements and the National Cancer Drugs Fund requirements
<a href="#">Pirfenidone for treating idiopathic pulmonary fibrosis (TA504)</a>	06/02/2018	<b>Pirfenidone (Esbriet®)</b> is recommended as an option for treating idiopathic pulmonary fibrosis in adults	x		28/03/2018	50	NHSE commissioned. Approved for addition to the formulary and to be used in line with NICE guidance and commissioning statements

<a href="#">Fulvestrant for untreated locally advanced or metastatic oestrogen-receptor positive breast cancer (TA503)</a>	31/01/2018	<b>Fulvestrant (Faslodex®)</b> is not recommended, within its marketing authorisation, for treating locally advanced or metastatic oestrogen-receptor positive breast cancer in postmenopausal women who have not had endocrine therapy before	x		28/03/2018	56	Not recommended
<a href="#">Ibrutinib for treating relapsed or refractory mantle cell lymphoma (TA502)</a>	31/01/2018	<b>Ibrutinib (Imbruvica®)</b> is recommended as an option for treating relapsed or refractory mantle cell lymphoma in adults	x		28/03/2018	56	NHSE commissioned. Approved for addition to the formulary and to be used in line with NICE guidance and commissioning statements
<a href="#">Intrabeam radiotherapy system for adjuvant treatment of early breast cancer (TA501)</a>	31/01/2018	<b>The Intrabeam radiotherapy system</b> is not recommended for routine commissioning for adjuvant treatment of early invasive breast cancer during breast-conserving surgical removal of the tumour	x		28/03/2018	56	Not recommended
<a href="#">Ceritinib for untreated ALK-positive non-small-cell lung cancer (TA500)</a>	24/01/2018	<b>Ceritinib (Zykadia®)</b> is recommended, within its marketing authorisation, as an option for untreated anaplastic lymphoma kinase (ALK)-positive advanced non-small-cell lung cancer in adults	x		28/03/2018	63	NHSE commissioned. Approved for addition to the formulary and to be used in line with NICE guidance and commissioning statements
<a href="#">Glecaprevir–pibrentasvir for treating chronic hepatitis C (TA499)</a>	24/01/2018	<b>Glecaprevir–pibrentasvir (Maviret®)</b> is recommended, within its marketing authorisation, as an option for treating chronic hepatitis C in adults	x		28/03/2018	63	NHSE commissioned. Approved for addition to the formulary and to be used in line with NICE guidance and commissioning statements
<a href="#">Lenvatinib with everolimus for previously treated advanced renal cell carcinoma (TA498)</a>	24/01/2018	<b>Lenvatinib (Kisplyx®) plus everolimus</b> is recommended as an option for treating advanced renal cell carcinoma in adults who have had 1 previous vascular endothelial growth factor (VEGF)-targeted therapy	x		28/03/2018	63	NHSE commissioned. Approved for addition to the formulary and to be used in line with NICE guidance and commissioning statements
<a href="#">Golimumab for treating non-radiographic axial spondyloarthritis (TA497)</a>	10/01/2018	<b>Golimumab (Simponi®)</b> is recommended, within its marketing authorisation, as an option for treating severe non-radiographic axial spondyloarthritis in adults whose disease has responded inadequately to, or who cannot tolerate, nonsteroidal anti-inflammatory drugs.	x		24/01/2018	14	CCG commissioned. Approved for addition to the formulary and to be used in line with NICE guidance and commissioning statements
<a href="#">Ribociclib with an aromatase inhibitor for previously untreated, hormone receptor-positive, HER2-negative, locally advanced or metastatic breast cancer (TA496)</a>	20/12/2017	<b>Ribociclib (Kisqali)</b> , with an aromatase inhibitor, is recommended within its marketing authorisation, as an option for treating hormone receptor-positive, human epidermal growth factor receptor 2-negative, locally advanced or metastatic breast cancer as initial endocrine-based therapy in adults	x		24/01/2018	35	NHSE commissioned. Approved for addition to the formulary and to be used in line with NICE guidance and commissioning statements
<a href="#">Palbociclib with an aromatase inhibitor for previously untreated, hormone receptor-positive, HER2-negative, locally advanced or metastatic breast cancer (TA495)</a>	20/12/2017	<b>Palbociclib (Ibrance)</b> , with an aromatase inhibitor, is recommended within its marketing authorisation, as an option for treating hormone receptor-positive, human epidermal growth factor receptor 2-negative, locally advanced or metastatic breast cancer as initial endocrine-based therapy in adults	x		24/01/2018	35	NHSE commissioned. Approved for addition to the formulary and to be used in line with NICE guidance and commissioning statements

<a href="#">Naltrexone–bupropion for managing overweight and obesity (TA494)</a>	12/12/2017	<b>Naltrexone–bupropion (Mysimba)</b> is not recommended within its marketing authorisation for managing overweight and obesity in adults alongside a reduced-calorie diet and increased physical activity	x		24/01/2018	43	Not recommended
<a href="#">Cladribine tablets for treating relapsing–remitting multiple sclerosis (TA493)</a>	06/12/2017	<b>Cladribine tablets (Mavenclad)</b> are recommended as an option for treating highly active multiple sclerosis in adults	x		24/01/2018	49	NHSE commissioned. Approved for addition to the formulary and to be used in line with NICE guidance and commissioning statements and the National Cancer Drugs Fund requirements
<a href="#">Atezolizumab for untreated locally advanced or metastatic urothelial cancer when cisplatin is unsuitable (TA492)</a>	06/12/2017	<b>Atezolizumab (Tecentriq)</b> is recommended for use within the Cancer Drugs Fund as an option for untreated locally advanced or metastatic urothelial carcinoma in adults, for whom cisplatin-based chemotherapy is unsuitable	x		24/01/2018	49	NHSE commissioned. Approved for addition to the formulary and to be used in line with NICE guidance and commissioning statements and the National Cancer Drugs Fund requirements
<a href="#">Ibrutinib for treating Waldenstrom's macroglobulinaemia (TA491)</a>	22/11/2017	<b>Ibrutinib (Imbruvica)</b> is recommended for use in the Cancer Drugs Fund as an option for treating Waldenstrom's macroglobulinaemia in adults who have had at least 1 prior therapy	x		24/01/2018	63	NHSE commissioned. Approved for addition to the formulary and to be used in line with NICE guidance and commissioning statements and the National Cancer Drugs Fund requirements
<a href="#">Nivolumab for treating squamous cell carcinoma of the head and neck after platinum-based chemotherapy (TA490)</a>	22/11/2017	<b>Nivolumab (Opdivo)</b> is recommended for use within the Cancer Drugs Fund as an option for treating squamous cell carcinoma of the head and neck in adults whose disease has progressed on platinum-based chemotherapy	x		24/01/2018	63	NHSE commissioned. Approved for addition to the formulary and to be used in line with NICE guidance and commissioning statements and the National Cancer Drugs Fund requirements
<a href="#">Vismodegib for treating basal cell carcinoma (TA489)</a>	22/11/2017	<b>Vismodegib (Erivedge)</b> is not recommended within its marketing authorisation for treating symptomatic metastatic basal cell carcinoma, or locally advanced basal cell carcinoma that is inappropriate for surgery or radiotherapy	x		24/01/2018	63	Not recommended
<a href="#">Regorafenib for previously treated unresectable or metastatic gastrointestinal stromal tumours (TA488)</a>	15/11/2017	<b>Regorafenib (Stivarga)</b> is recommended as an option for treating unresectable or metastatic gastrointestinal stromal tumours in adults whose disease has progressed on, or who are intolerant to, prior treatment with imatinib and sunitinib	x		24/01/2018	70	NHSE commissioned. Approved for addition to the formulary and to be used in line with NICE guidance and commissioning statements and the National Cancer Drugs Fund requirements
<a href="#">Venetoclax for treating chronic lymphocytic leukaemia (TA487)</a>	01/10/2017	<b>Venetoclax (Venclyxto®)</b> 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	NHSE commissioned. Approved for addition to the formulary and to be used in line with NICE guidance and commissioning statements and the National Cancer Drugs Fund requirements
<a href="#">Aflibercept for treating choroidal neovascularisation (TA486)</a>	01/10/2017	<b>Aflibercept (Eylea®)</b> 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	CCG commissioned. Approved for addition to the formulary and to be used in line with NICE guidance and commissioning statements
<a href="#">Sarilumab for moderate to severe rheumatoid arthritis (TA485)</a>	01/10/2017	<b>Sarilumab (Kevzara®)</b> 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	CCG commissioned. Approved for addition to the formulary and to be used in line with NICE guidance and commissioning statements

<a href="#">Nivolumab for previously treated non-squamous non-small-cell lung cancer (TA484)</a>	01/10/2017	<b>Nivolumab (Opdivo®)</b> 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	NHSE commissioned. Approved for addition to the formulary and to be used in line with NICE guidance and commissioning statements and the National Cancer Drugs Fund requirements
<a href="#">Nivolumab for previously treated squamous non-small-cell lung cancer (TA483)</a>	01/10/2017	<b>Nivolumab (Opdivo®)</b> 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	NHSE commissioned. Approved for addition to the formulary and to be used in line with NICE guidance and commissioning statements and the National Cancer Drugs Fund requirements
<a href="#">Immunosuppressive therapy for kidney transplant in children and young people (TA482)</a>	01/10/2017	Immunosuppressive therapy for kidney transplant in children and young people	x		29/11/2017	59	NHSE commissioned. Approved for addition to the formulary and to be used in line with NICE guidance and commissioning statements
<a href="#">Immunosuppressive therapy for kidney transplant in adults (TA481)</a>	01/10/2017	Immunosuppressive therapy for kidney transplant in adults	x		29/11/2017	59	NHSE commissioned. Approved for addition to the formulary and to be used in line with NICE guidance and commissioning statements
<a href="#">Tofacitinib for moderate to severe rheumatoid arthritis (TA480)</a>	01/10/2017	<b>Tofacitinib (Xeljanz®)</b> 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	CCG commissioned. Approved for addition to the formulary and to be used in line with NICE guidance and commissioning statements
<a href="#">Reslizumab for treating severe eosinophilic asthma (TA479)</a>	01/10/2017	<b>Reslizumab (Cinqaero®)</b> 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	NHSE commissioned. Approved for addition to the formulary and to be used in line with NICE guidance and commissioning statements
<a href="#">Brentuximab vedotin for treating relapsed or refractory systemic anaplastic large cell lymphoma (TA478)</a>	01/10/2017	<b>Brentuximab vedotin (Adcetris®)</b> is recommended as an option for treating relapsed or refractory systemic anaplastic large cell lymphoma in adults	x		29/11/2017	59	NHSE commissioned. Approved for addition to the formulary and to be used in line with NICE guidance and commissioning statements and the National Cancer Drugs Fund requirements
<a href="#">Autologous chondrocyte implantation for treating symptomatic articular cartilage defects of the knee (TA477)</a>	01/10/2017	<b>Autologous chondrocyte implantation (ACI) ChondroCelect®</b> 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
<a href="#">Paclitaxel as albumin-bound nanoparticles with gemcitabine for untreated metastatic pancreatic cancer (TA476)</a>	06/09/2017	<b>Paclitaxel as albumin-bound nanoparticles (nab-paclitaxel) with gemcitabine</b> 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
<a href="#">Dimethyl fumarate for treating moderate to severe plaque psoriasis (TA475)</a>	06/09/2017	<b>Dimethyl fumarate</b> is recommended as an option for treating plaque psoriasis in adults	x		29/11/2017	84	Confirmed place in therapy with dermatologists. CCG commissioned. Approved for addition to the formulary and to be used in line with NICE guidance and commissioning statements.
<a href="#">Sorafenib for treating advanced hepatocellular carcinoma (TA474)</a>	06/09/2017	<b>Sorafenib</b> 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

<a href="#">Cetuximab for treating recurrent or metastatic squamous cell cancer of the head and neck (TA473)</a>	31/08/2017	<b>Cetuximab in combination with platinum-based chemotherapy</b> 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.
<a href="#">Obinutuzumab with bendamustine for treating follicular lymphoma refractory to rituximab (TA472)</a>	30/08/2017	<b>Obinutuzumab in combination with bendamustine followed by obinutuzumab maintenance</b> 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
<a href="#">Eluxadoline for treating irritable bowel syndrome with diarrhoea (TA471)</a>	30/08/2017	<b>Eluxadoline</b> 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.
<a href="#">Ofatumumab with chemotherapy for treating chronic lymphocytic leukaemia (terminated appraisal) (TA470)</a>	23/08/2017	<b>Ofatumumab with chemotherapy</b> - 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
<a href="#">Idelalisib with ofatumumab for treating chronic lymphocytic leukaemia (terminated appraisal) (TA469)</a>	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
<a href="#">Methylnaltrexone bromide for treating opioid-induced constipation (terminated appraisal) (TA468)</a>	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
<a href="#">Holoclar for treating limbal stem cell deficiency after eye burns (TA467)</a>	16/08/2017	<b>Holoclar (ex vivo expanded autologous human corneal epithelial cells containing stem cells)</b> 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
<a href="#">Baricitinib for moderate to severe rheumatoid arthritis (TA466)</a>	09/08/2017	<b>Baricitinib, with methotrexate</b> 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
<a href="#">Olaratumab in combination with doxorubicin for treating advanced soft tissue sarcoma (TA465)</a>	09/08/2017	<b>Olaratumab, in combination with doxorubicin,</b> 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

<a href="#">Bisphosphonates for treating osteoporosis (TA464)</a>	09/08/2017	<b>Oral bisphosphonates</b> (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
<a href="#">Cabozantinib for previously treated advanced renal cell carcinoma (TA463)</a>	09/08/2017	<b>Cabozantinib</b> 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
<a href="#">Nivolumab for treating relapsed or refractory classical Hodgkin lymphoma (TA462)</a>	26/07/2017	<b>Nivolumab</b> 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
<a href="#">Roflumilast for treating chronic obstructive pulmonary disease (TA461)</a>	26/07/2017	<b>Roflumilast</b> , 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
<a href="#">Adalimumab and dexamethasone for treating non-infectious uveitis (TA460)</a>	26/07/2017	<b>Adalimumab</b> 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
<a href="#">Collagenase clostridium histolyticum for treating Dupuytren's contracture (TA459)</a>	26/07/2017	<b>Collagenase clostridium histolyticum</b> 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
<a href="#">Trastuzumab emtansine for treating HER2-positive advanced breast cancer after trastuzumab and a taxane (TA458)</a>	19/07/2017	<b>Trastuzumab emtansine</b> 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

<a href="#">Carfilzomib for previously treated multiple myeloma (TA457)</a>	19/07/2017	<b>Carfilzomib in combination with dexamethasone</b> 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
<a href="#">Ustekinumab for moderately to severely active Crohn's disease after previous treatment (TA456)</a>	12/07/2017	<b>Ustekinumab</b> 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
<a href="#">Adalimumab, etanercept and ustekinumab for treating plaque psoriasis in children and young people (TA455)</a>	12/07/2017	<b>Adalimumab, Etanercept and Ustekinumab</b> 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
<a href="#">Daratumumab with lenalidomide and dexamethasone for treating relapsed or refractory multiple myeloma (terminated appraisal) (TA454)</a>	05/07/2017	<b>Daratumumab</b> - 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
<a href="#">Bortezomib for treating multiple myeloma after second or subsequent relapse (terminated appraisal) (TA453)</a>	05/07/2017	<b>Bortezomib</b> - 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
<a href="#">Ibrutinib for untreated chronic lymphocytic leukaemia without a 17p deletion or TP53 mutation (terminated appraisal) (TA452)</a>	05/07/2017	<b>Ibrutinib</b> - 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
<a href="#">Ponatinib for treating chronic myeloid leukaemia and acute lymphoblastic leukaemia (TA451)</a>	28/06/2017	<b>Ponatinib (Iclusig)</b> 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
<a href="#">Blinatumomab for previously treated Philadelphia-chromosome-negative acute lymphoblastic leukaemia (TA450)</a>	28/06/2017	<b>Blinatumomab (Blinicyto)</b> 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

<a href="#">Everolimus and sunitinib for treating unresectable or metastatic neuroendocrine tumours in people with progressive disease (TA449)</a>	28/06/2017	<b>Everolimus (Afinitor) and Sunitinib (Sutent)</b> 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
<a href="#">Etelcalcetide for treating secondary hyperparathyroidism (TA448)</a>	28/06/2017	<b>Etelcalcetide (Parsabiv)</b> 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
<a href="#">Pembrolizumab for untreated PD-L1-positive metastatic non-small-cell lung cancer (TA447)</a>	28/06/2017	<b>Pembrolizumab (Keytruda)</b> 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
<a href="#">Brentuximab vedotin for treating CD30-positive Hodgkin lymphoma (TA446)</a>	28/06/2017	<b>Brentuximab vedotin (Adcetris)</b> 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
<a href="#">Certolizumab pegol and secukinumab for treating active psoriatic arthritis after inadequate response to DMARDs (TA445)</a>	24/05/2017	<b>Certolizumab pegol (Cimzia) alone, or in combination with methotrexate</b> 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.
<a href="#">Afinib for treating advanced squamous non-small-cell lung cancer after platinum-based chemotherapy (terminated appraisal) (TA444)</a>	22/03/2017	<b>Tenofovir alafenamide</b> - was unable to make a recommendation about the use in the NHS of afinib 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
<a href="#">Obeticholic acid for treating primary biliary cholangitis (TA443)</a>	26/04/2017	<b>Obeticholic acid (Ocaliva)</b> 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
<a href="#">Ixekizumab for treating moderate to severe plaque psoriasis (TA442)</a>	26/04/2017	<b>Ixekizumab (Taltz)</b> 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.
<a href="#">Daclizumab for treating relapsing–remitting multiple sclerosis (TA441)</a>	26/04/2017	<b>Daclizumab (Zinbryta)</b> 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



<a href="#">Pegylated liposomal irinotecan for treating pancreatic cancer after gemcitabine (TA440)</a>	26/04/2017	<b>Pegylated liposomal irinotecan (Onivyde)</b> , 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
			70	0			
			% "Yes"	% "N/A"	-	<b>Average implement time (days)</b>	
<b>Adherence statistics for 2017-18</b>			100%	0%		<b>47</b>	