Milton Keynes Health Economy

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

This spreadsheet is updated monthly and enables self-audit of a medicines formulary for adherence to current NICE Technology Appraisals. January 2019 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) | |
| 2018/2019 Pembrolizumab with pemetrexed and platinum chemotherapy for untreated, metastatic, non-squamous non- small-cell lung cancer (TA557) | | Pembrolizumab, with pemetrexed and platinum chemotherapy is recommended for use within the Cancer Drugs Fund, as an option for untreated, metastatic, non-squamous non-small- cell lung cancer (See criteria via link) | x | | 23/01/2019 | | NHSE / CDF commissioned. Approved for addition to the formulary and to be used in line with NICE TA557 and commissioning statements | |
| Darvadstrocel for treating complex perianal fistulas in Crohn's disease (TA556) | 09/01/2019 | Darvadstrocel is not recommended, within its marketing authorisation, for previously treated complex perianal fistulas in adults with non- active or mildly active luminal Crohn's disease. | x | | 23/01/2019 | | Not recommended | |
| Regorafenib for previously treated advanced hepatocellular carcinoma (TA555) | 09/01/2019 | Regorafenib is recommended as an option for treating advanced unresectable hepatocellular carcinoma in adults who have had sorafenib (SEE CRITERIA VIA LINK) | x | | 23/01/2019 | | NHSE / CDF commissioned. Approved for addition to the formulary and to be used in line with NICE TA555 and commissioning statements | |
| Tisagenlecleucel for treating relapsed or refractory B-cell acute lymphoblastic leukaemia in people aged up to 25 years (TA554) | | Tisagenlecleucel therapy is recommended for use within the Cancer Drugs Fund as an option for treating relapsed or refractory B-cell acute lymphoblastic leukaemia in people aged up to 25 years | x | | 23/01/2019 | | NHSE / CDF commissioned. Approved for addition to the formulary and to be used in line with NICE TA554 and commissioning statements | |
| Pembrolizumab for adjuvant treatment of resected melanoma with high risk of recurrence (TA553) | 19/12/2018 | Pembrolizumab is recommended for use within the Cancer Drugs Fund as an option for the adjuvant treatment of stage III melanoma with lymph node involvement in adults who have had complete resection | x | | 23/01/2019 | | NHSE commissioned. Approved for addition to the formulary and to be used in line with NICE TA553 and commissioning statements | |

| Liposomal_ cytarabine-daunorubicin for_ untreated acute myeloid_ leukaemia (TA552) | 19/12/2018 | Liposomal cytarabine–daunorubicin is recommended, within its marketing authorisation, as an option for untreated therapy-related acute myeloid leukaemia or acute myeloid leukaemia with myelodysplasia- | x | 23/01/2019 | NHSE commissioned. Approved for addition to the formulary and to be used in line with NICE TA552 and commissioning statements |
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| Lenvatinib for untreated advanced hepatocellular carcinoma (TA551) | 19/12/2018 | Lenvatinib is recommended as an option for untreated, advanced, unresectable hepatocellular carcinoma in adults (see criteria via link) | x | 23/01/2019 | NHSE commissioned. Approved for addition to the formulary and to be used in line with NICE TA551 and commissioning statements |
| Vandetanib for treating medullary thyroid cancer (TA550) | 12/12/2018 | Vandetanib is not recommended, within its marketing authorisation, for treating aggressive and symptomatic medullary thyroid cancer in adults with unresectable, locally advanced or metastatic disease. | х | 23/01/2019 | Not recommended |
| Denosumab for preventing skeletal-related events in multiple myeloma (terminated appraisal) (TA549) | 05/12/2018 | NICE is unable to make a recommendation about the use in the NHS of denosumab for preventing skeletal-related events in multiple myeloma because no evidence submission was received from Amgen | x | 23/01/2019 | Terminated Appraisal |
| Decitabine for untreated acute myeloid leukaemia (terminated appraisal) (TA548) | 05/12/2018 | NICE is unable to make a recommendation about the use in the NHS of decitabine for untreated acute myeloid leukaemia because no evidence submission was received from Janssen. | x | 23/01/2019 | Terminated Appraisal |
| Tofacitinib for moderately to severely active ulcerative colitis (TA547) | 28/11/2018 | Tofacitinib is recommended, within its marketing authorisation, as an option for treating moderately to severely active ulcerative colitis in adults when conventional therapy or a biological agent cannot be tolerated or the disease has responded inadequately or lost response to treatment. | x | 23/01/2019 | CCG commissioned. Approved for addition to the formulary and to be used in line with NICE TA547 and commissioning statements; Blueteq form required |
| Padeliporfin for untreated localised prostate cancer (TA546) | 21/11/2018 | Padeliporfin is not recommended, within its marketing authorisation, for untreated, unilateral, low-risk prostate cancer in adults | х | 23/01/2019 | Not recommended |

| Gemtuzumab ozogamicin for untreated acute myeloid leukaemia (TA545) | 14/11/2018 | Gemtuzumab ozogamicin, with daunorubicin and cytarabine, is recommended as an option for untreated de novo CD33-positive acute myeloid leukaemia (AML), except acute promyelocytic leukaemia, in people 15 years and over | Х | 23/01/2019 | | NHSE commissioned. Approved for addition to the formulary and to be used in line with NICE TA545 and commissioning statements |
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| Dabrafenib with trametinib for adjuvant treatment of resected BRAF V600 mutation-positive melanoma (TA544) | 17/10/2018 | Dabrafenib with trametinib is recommended, within its marketing authorisation, as an option for the adjuvant treatment of resected stage III BRAF V600 mutation-positive melanoma in adults | х | 28/11/2019 | | NHSE commissioned. Approved for addition to the formulary and to be used in line with NICE TA544 and commissioning statements |
| Tofacitinib for treating active psoriatic arthritis after inadequate response to DMARDs (TA543) | 03/10/2018 | Tofacitinib, with methotrexate, is recommended as an option for treating active psoriatic arthritis in adults, only if: (see criteria via link) | x | 28/11/2018 | 56 | CCG commissioned. Approved for addition to the formulary and to be used in line with NICE TA543 and commissioning statements |
| Cabozantinib for untreated advanced renal cell carcinoma (TA542) | 03/10/2018 | Cabozantinib is recommended, within its marketing authorisation, for adults with untreated advanced renal cell carcinoma that is intermediate- or poor-risk as defined in the International Metastatic Renal Cell Carcinoma | х | 28/11/2018 | 56 | NHSE commissioned. Approved for addition to the formulary and to be used in line with NICE TA542 and commissioning statements |
| Inotuzumab ozogamicin for treating relapsed or refractory B-cell acute lymphoblastic leukaemia (TA541) | 19/09/2018 | Inotuzumab ozogamicin is recommended, within its marketing authorisation, as an option for treating relapsed or refractory CD22-positive B- cell precursor acute lymphoblastic leukaemia in adults. | x | 28/11/2018 | | NHSE commissioned. Approved for addition to the formulary and to be used in line with NICE TA541 and commissioning statements |
| Pembrolizumab for treating relapsed or refractory classical Hodgkin lymphoma (TA540) | 03/09/2018 | Pembrolizumab is not recommended for treating relapsed or refractory classical Hodgkin lymphoma in adults who have had autologous stem cell transplant and brentuximab vedotin | x | 26/09/2018 | 23 | NHSE commissioned. Approved for addition to the formulary and to be used in line with NICE TA540 and commissioning statements |
| Lutetium (177Lu) oxodotreotide for treating unresectable or metastatic neuroendocrine tumours (TA539) | 29/08/2018 | Lutetium (177Lu) oxodotreotide is recommended, within its marketing authorisation, as an option for treating unresectable or metastatic, progressive, well- differentiated (grade 1 or grade 2), somatostatin receptor-positive gastroenteropancreatic neuroendocrine tumours (NETs) in adults | x | 26/09/2018 | 28 | NHSE commissioned. Approved for addition to the formulary and to be used in line with NICE TA539 and commissioning statements |

| Dinutuximab beta for treating neuroblastoma (TA538) | 22/08/2018 | Dinutuximab beta is recommended as an option for treating high-risk neuroblastoma in people aged 12 months and over whose disease has at least partially responded to induction chemotherapy, followed by myeloablative therapy and stem cell transplant | x | 26/09/2018 | 35 | NHSE commissioned. Approved for addition to the formulary and to be used in line with NICE TA538 and commissioning statements |
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| Ixekizumab for treating active psoriatic arthritis after inadequate response to DMARDs (TA537) | 08/08/2018 | Ixekizumab alone, or with methotrexate, is recommended as an option for treating active psoriatic arthritis in adults | х | 26/09/2018 | 49 | CCG commissioned. Approved for addition to the formulary and to be used in line with NICE TA537 and commissioning statements |
| Alectinib for untreated ALK- positive advanced non-small-cell lung cancer (TA536) | 08/08/2018 | Alectinib is recommended, within its marketing authorisation, as an option for untreated anaplastic lymphoma kinase (ALK)-positive advanced non-small-cell lung cancer (NSCLC) in adults | x | 26/09/2018 | 49 | NHSE commissioned. Approved for addition to the formulary and to be used in line with NICE guidance and commissioning statements |
| Lenvatinib and sorafenib for treating differentiated thyroid cancer after radioactive iodine (TA535) | 08/08/2018 | Lenvatinib and sorafenib are recommended as options for treating progressive, locally advanced or metastatic differentiated thyroid cancer (papillary, follicular or Hürthle cell) in adults whose disease does not respond to radioactive iodine | x | 26/09/2018 | 49 | NHSE commissioned. Approved for addition to the formulary and to be used in line with NICE guidance and commissioning statements |
| Dupilumab for treating moderate to severe atopic dermatitis (TA534) | 01/08/2018 | Dupilumab is recommended as an option for treating moderate to severe atopic dermatitis in adults | x | 26/09/2018 | 56 | CCG commissioned. Approved for addition to the formulary and to be used in line with NICE guidance and commissioning statements |
| Ocrelizumab for treating relapsing-remitting multiple sclerosis (TA533) | 25/07/2018 | Ocrelizumab is recommended as an option for treating relapsing-remitting multiple sclerosis in adults with active disease defined by clinical or imaging features | x | 26/09/2018 | 63 | NHSE commissioned. Approved for addition to the formulary and to be used in line with NICE guidance and commissioning statements |
| <u>Cenegermin for treating</u> neurotrophic keratitis (TA532) | 18/07/2018 | Cenegermin is not recommended, within its marketing authorisation, for treating moderate or severe neurotrophic keratitis in adults | x | 26/09/2018 | 70 | Not recommended |
| Pembrolizumab for untreated PD- L1-positive metastatic non-small- cell lung cancer (TA531) | 18/07/2018 | Pembrolizumab is recommended as an option for untreated PD-L1-positive metastatic non- small-cell lung cancer (NSCLC) in adults whose tumours express PD-L1 (with at least a 50% tumour proportion score) and have no epidermal growth factor receptor- or anaplastic lymphoma kinase-positive mutations. | x | 26/09/2018 | 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 |
| Nivolumab for treating locally advanced unresectable or metastatic urothelial cancer after platinum-containing chemotherapy (TA530) | 04/07/2018 | Nivolumab is not recommended, within its marketing authorisation, for treating locally advanced unresectable or metastatic urothelial carcinoma in adults who have had platinum- containing therapy. | x | 25/07/2018 | 21 | Not recommended |
| Crizotinib for treating ROS1- positive advanced non-small-cell lung cancer (TA529) | 04/07/2018 | | х | 25/07/2018 | 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 |

| Niraparib for maintenance treatment of relapsed, platinum- sensitive ovarian, fallopian tube and peritoneal cancer (TA528) Beta interferons and glatiramer acetate for treating multiple sclerosis (TA527) | 04/07/2018 | Niraparib is recommended for use within the Cancer Drugs Fund as an option for treating relapsed, platinum-sensitive high-grade serous epithelial ovarian, fallopian tube or primary peritoneal cancer that has responded to the most recent course of platinum-based Interferon beta-1a is recommended as an option for treating multiple sclerosis | x | 25/07/2018 | | 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 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 |
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| Arsenic trioxide for treating acute promyelocytic leukaemia (TA526) | 13/06/2018 | Arsenic trioxide is recommended, within its marketing authorisation, as an option for inducing remission and consolidation in acute promyelocytic leukaemia (characterised by the presence of the t[15;17] translocation or the PML/RAR-alpha gene) in adults | x | 25/07/2018 | 42 | 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 |
| Atezolizumab for treating locally advanced or metastatic urothelial carcinoma after platinum- containing chemotherapy (TA525) | 13/06/2018 | Atezolizumab is recommended as an option for treating locally advanced or metastatic urothelial carcinoma in adults who have had platinum-containing chemotherapy | x | 25/07/2018 | 42 | 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 |
| Brentuximab vedotin for treating CD30-positive Hodgkin lymphoma (TA524) | 13/06/2018 | Brentuximab vedotin is recommended as an option for treating CD30-positive Hodgkin lymphoma in adults with relapsed or refractory disease | x | 25/07/2018 | 42 | 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 |
| Midostaurin for untreated acute myeloid leukaemia (TA523) | 13/06/2018 | Midostaurin is recommended, within its marketing authorisation, as an option in adults for treating newly diagnosed acute FLT3- mutation-positive myeloid leukaemia with | x | 25/07/2018 | 42 | 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 |
| Pembrolizumab for untreated PD- L1-positive locally advanced or metastatic urothelial cancer when cisplatin is unsuitable (TA522) | 13/06/2018 | Pembrolizumab is recommended for use within the Cancer Drugs Fund as an option for untreated locally advanced or metastatic urothelial carcinoma in adults when cisplatin- containing chemotherapy is unsuitable | x | 25/07/2018 | 42 | 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 |
| Guselkumab for treating moderate to severe plaque psoriasis (TA521) | 13/06/2018 | Guselkumab is recommended as an option for treating plaque psoriasis in adults | x | 25/07/2018 | 42 | CCG commissioned. Approved for addition to the formulary and to be used in line with NICE guidance and commissioning statements |
| Atezolizumab for treating locally advanced or metastatic non-small- cell lung cancer after chemotherapy (TA520) | 16/04/2018 | Atezolizumab is recommended as an option for treating locally advanced or metastatic non-small-cell lung cancer (NSCLC) in adults who have had chemotherapy (and targeted treatment if they have an EGFR- or ALK-positive tumour), | x | 23/05/2018 | 37 | 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 |
| Pembrolizumab for treating locally advanced or metastatic urothelial carcinoma after platinum-containing chemotherapy (TA519) | 25/04/2018 | Pembrolizumab is recommended for use within the Cancer Drugs Fund as an option for treating locally advanced or metastatic urothelial carcinoma in adults who have had platinum- containing chemotherapy | x | 23/05/2018 | 28 | 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 |

| Tocilizumab for treating giant cell arteritis (TA518) | | Tocilizumab, when used with a tapering course of glucocorticoids (and when used alone after glucocorticoids), is recommended as an option for treating giant cell arteritis in adults | x | | 23/05/2018 | | 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 |
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| Avelumab for treating metastatic Merkel cell carcinoma (TA517) | 11/04/2018 | Avelumab is recommended as an option for treating metastatic Merkel cell carcinoma in adults, only if they have had 1 or more lines of chemotherapy for metastatic disease. | x | | 23/05/2018 | | 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 |
| | | | % "Yes" | % "N/A" | - | Average implement time (days) | |
| Adherence statistics for 2018-19 | | | 100% | 0% | | 41 | |