Formulary Group work programme

The decisions of the Formulary Group meeting (Tuesday 18 June 2024) will be published by 03 July 2024.

Publication schedule

This information is updated monthly.

Formulary Group work programme

The Group meets monthly on the third Tuesday of each month (excluding July and December). The following list includes medicines that will be scheduled for review in the next three to four months. It is indicative and may be subject to change.

Medicine and indication	Meeting date
acarbose 50mg, 100mg tablets	
Formulary review	
Indication: recommended for the treatment of type 2 diabetes (non-insulin	
dependent) in patients inadequately controlled on diet alone, or on diet and	Jun 2024
(i) metformin and / or (ii) a sulphonylurea.	
atovaquone 750mg/5mL oral suspension	
FG1 451/22	
Indication: (off-label) for the prevention of Pneumocystis carinii	
pneumonia.	ТВС
Restriction: second-line option for patients where co-trimoxazole is not	TDC
effective, not tolerated or contraindicated.	
clonidine 25micrograms, 100micrograms tablets	
Formulary review	
Indication: off-label for children and adolescents 6 years and older for the	
management of ADHD in those unresponsive to or unable to tolerate	Jun 2024
treatment with stimulants, atomoxetine or guanfacine.	
dostarlimab 500mg concentrate for solution for infusion (Jemperli®)	
SMC 2635	
Indication: in combination with platinum-containing chemotherapy for the	
treatment of adult patients with mismatch repair deficient	Sep 2024
(dMMR)/microsatellite instability-high (MSI-H) primary advanced or	3ep 2024
recurrent endometrial cancer and who are candidates for systemic therapy.	
dulaglutide 0.75mg, 1.5mg, 3mg, 4.5mg solution for injection in pre-filled pen (Trulicity®)	
Formulary review 1110/15	
Indication: in adults with type 2 diabetes mellitus to improve glycaemic	
control as add-on therapy in combination with other glucose-lowering	
medicinal products including insulin, when these, together with diet and	
exercise, do not provide adequate glycaemic control.	Jun 2024
SMC restriction: as part of a triple therapy in patients with inadequate	
glycaemic control on two oral anti-diabetic drugs, as an alternative	
glucagon-like peptide 1 (GLP-1) agonist option.	

Medicine and indication	Meeting date
ertugliflozin L-pyroglutamic acid 5mg, 15mg tablets	
Formulary review 2102	
Indication: in adults aged 18 years and older with type 2 diabetes mellitus	
as an adjunct to diet and exercise to improve glycaemic control:	
As monotherapy in patients for whom the use of metformin is considered	
inappropriate due to intolerance or contraindications.	
In addition to other medicinal products for the treatment of diabetes.	Jun 2024
SMC restriction: ertugliflozin is accepted for use as monotherapy and as	
add-on therapy. When used as monotherapy it is restricted to patients who	
would otherwise receive a dipeptidyl peptidase-4 inhibitor and in whom a	
sulphonylurea or pioglitazone is not appropriate.	
exenatide 2mg powder and solvent for prolonged-release suspension for	
injection (Bydureon®)	
Formulary review 748/11	
Indication: treatment of type 2 diabetes mellitus in combination with:	
- metformin	
- sulphonylurea	
- thiazolidinedione	
- metformin and sulphonylurea	Jun 2024
- metformin and thiazolidinedione in adults who have not achieved	3411 202 1
adequate glycaemic control on maximally tolerated doses of these oral	
therapies.	
SMC restriction: exenatide once weekly is restricted to use as a third line	
treatment option.	
foslevodopa-foscarbidopa 240mg/mL + 12mg/mL solution for infusion	
(Produodopa®)	
SMC 2574	
Indication: treatment of advanced levodopa-responsive Parkinson's disease	
with severe motor fluctuations and hyperkinesia or dyskinesia when	
available combinations of Parkinson medicinal products have not given	Jun 2024
satisfactory results.	-
SMC restriction: for use in patients not eligible for deep brain stimulation	
(DBS).	
glofitamab 2.5mg, 10mg concentrate for solution for infusion (Columvi®)	
SMC 2614	
Indication: as monotherapy for the treatment of adult patients with	
relapsed or refractory (R/R) diffuse large B-cell lymphoma (DLBCL), after	TBC
two or more lines of systemic therapy.	

Medicine and indication	Meeting date
GoResp® Digihaler® 160micrograms/4.5micrograms, 320micrograms/9micrograms budesonide / formoterol fumarate dihydrate	
Formulary review	
Indication: GoResp Digihaler is indicated in adults 18 years of age and older only. Asthma GoResp Digihaler is indicated in the regular treatment of asthma, where use of a combination (inhaled corticosteroid and long-acting β2 adrenoceptor agonist) is appropriate: -in patients not adequately controlled with inhaled corticosteroids and "as needed" inhaled short-acting β2 adrenoceptor agonists. or -in patients already adequately controlled on both inhaled corticosteroids and long-acting β2 adrenoceptor agonists.	Jun 2024
COPD Symptomatic treatment of patients with COPD with forced expiratory volume in 1 second (FEV1) < 70% predicted normal (post bronchodilator) and a history of repeated exacerbations, who have significant symptoms despite regular therapy with long-acting bronchodilators.	
ivosidenib 250mg film-coated tablet (Tibsovo®) SMC 2615	
Indication: in combination with azacitidine for the treatment of adult patients with newly diagnosed acute myeloid leukaemia (AML) with an isocitrate dehydrogenase-1 (IDH1) R132 mutation who are not eligible to receive standard induction chemotherapy.	Aug 2024
liraglutide 6mg/mL solution for injection in pre-filled pen (Saxenda®)	
SMC 2455	
Indication: as an adjunct to a reduced-calorie diet and increased physical activity for weight management in adult patients with an initial Body Mass Index (BMI) of: 1) ≥30kg/m² (obese), or 2) ≥27kg/m² to <30kg/m² (overweight) in the presence of at least one weight-related comorbidity such as dysglycaemia (prediabetes or type 2 diabetes mellitus), hypertension, dyslipidaemia or obstructive sleep apnoea. SMC restriction: BMI ≥35kg/m²* (obesity class II and above) with: - Non-diabetic hyperglycaemia (prediabetes) at high risk of type 2 diabetes which is defined as having either:o Fasting plasma glucose level of 5.5 to 6.9mmol/L or o HbA1c of 6.0 to 6.4% (42 to 47mmol/mol), and - High risk of cardiovascular disease (CVD): o Total cholesterol >5mmol/L, oro High-density lipoprotein (HDL) <1.0mmol/L for men and <1.3mmol/L for women, or o Systolic blood pressure (SBP) >140mmHg.Patients should be treated in a specialist weight management service. *a lower BMI cut-off may be more appropriate for members of minority	Jun 2024

Medicine and indication	Meeting date
ethnic groups known to be at equivalent risk of the consequences of obesity at a lower BMI than the white population.	
mavacamten 2.5mg, 5mg, 10mg, 15mg hard capsules (Camzyos®)	
SMC 2618	
Indication: treatment of symptomatic (New York Heart Association, NYHA, class II to III) obstructive hypertrophic cardiomyopathy (oHCM) in adult patients.	Aug 2024
mirikizumab 100mg, 300mg solution for injection in pre-filled pen and concentrate for solution for infusion (Omvoh®)	
SMC 2650	
Indication: for the treatment of adult patients with moderately to severely active ulcerative colitis who have had an inadequate response with, lost response to, or were intolerant to either conventional therapy or a biologic treatment.	Aug 2024
momelotinib 100mg, 150mg, 200mg film coated tablet (Omjjara®)	
SMC 2636	
Indication: for the treatment of disease-related splenomegaly or symptoms in adult patients with moderate to severe anaemia who have primary myelofibrosis, post polycythaemia vera myelofibrosis or post essential thrombocythaemia myelofibrosis and who are Janus Associated Kinase (JAK) inhibitor naïve or have been treated with ruxolitinib.	ТВС
repaglinide 0.5mg, 1mg, 2mg tablets	
Formulary review	
Indication: adults with type 2 diabetes mellitus whose hyperglycaemia can no longer be controlled satisfactorily by diet, weight reduction and exercise. Repaglinide is also indicated in combination with metformin in adults with type 2 diabetes mellitus who are not satisfactorily controlled on metformin alone.	Jun 2024
ribociclib 200mg film-coated tablets (Kisqali®)	
SMC 2198	
Indication: for the treatment of women with hormone receptor (HR)- positive, human epidermal growth factor receptor 2 (HER2)-negative locally advanced or metastatic breast cancer in combination with fulvestrant* as initial endocrine-based therapy, or in women who have received prior endocrine therapy. SMC restriction: women who have relapsed on or within 12 months of completing (neo) adjuvant endocrine therapy, or those who have progressed on first-line endocrine-based therapy for advanced breast cancer.	TBC
ritlecitinib tosylate 50mg hard capsules (Litfulo®)	
SMC 2610 Indication: for the treatment of severe alopecia areata in adults and adolescents 12 years of age and older.	Sep 2024

Medicine and indication	Meeting date
secukinumab 150mg, 300mg solution for injection in pre-filled pen or pre-filled syringe (Cosentyx®)	
SMC 2592	
Indication: for the treatment of active moderate to severe hidradenitis suppurativa (HS) (acne inversa) in adults with an inadequate response to conventional systemic HS therapy. SMC restriction: for use in adult patients with active moderate to severe HS for whom adalimumab is contraindicated or otherwise unsuitable, including those who have failed to respond or have lost response to prior adalimumab treatment.	Sep 2024
selpercatinib 40mg, 80mg hard capsules (Retsevmo®)	
SMC 2573	
Indication: monotherapy for the treatment of adults with advanced rearranged during transfection (RET) fusion-positive non-small cell lung cancer (NSCLC) not previously treated with a RET inhibitor. SMC restriction: for use in treatment-naïve patients who have not previously received a RET-inhibitor or any other systemic treatments for their advanced stage of disease.	Jun 2024
semaglutide 0.25mg, 0.5mg, 1mg, 2mg solution for injection in pre-filled pen (Ozempic®)	
Formulary review 2092	
Indication: the treatment of adults with insufficiently controlled type 2 diabetes mellitus (T2DM) as an adjunct to diet and exercise: - as monotherapy when metformin is considered inappropriate due to intolerance or contraindications - in addition to other medicinal products for the treatment of diabetes. SMC restriction: In addition to other oral anti-diabetic medicines, or as an add-on to basal insulin, as an alternative glucagon-like peptide-1 receptor agonist option.	Jun 2024
semaglutide 3mg, 7mg and 14mg tablets (Rybelsus®)	
SMC 2287	
Indication: for the treatment of adults with insufficiently controlled type 2 diabetes mellitus to improve glycaemic control as an adjunct to diet and exercise - as monotherapy when metformin is considered inappropriate due to intolerance or contraindications - in combination with other medicinal products for the treatment of diabetes. SMC restriction: In addition to other oral anti-diabetic medicines, or as an add-on to basal insulin, as an alternative glucagon-like peptide-1 receptor agonist option.	Jun 2024

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Medicine and indication	Meeting date
sodium zirconium cyclosilicate 5g and 10g powder for oral suspension (Lokelma®)	
SMC 2288	
Indication: for the treatment of hyperkalaemia in adult patients. SMC restriction: patients with hyperkalaemia (defined as a serum potassium of >6.0mmol/L) with chronic kidney disease (CKD) stage 3b to 5 and/or heart failure who would otherwise need to down-titrate or discontinue their renin-angiotensin-aldosterone system inhibitor (RAASi) therapy to maintain a clinically acceptable serum potassium level (normokalaemia)	Aug 2024
tirzepatide 2.5mg, 5mg, 7.5mg, 10mg, 12.5mg, 15mg solution for injection	
in pre-filled pen (Mounjaro®)	
SMC 2633 Indication: for the treatment of adults with insufficiently controlled type 2 diabetes mellitus as an adjunct to diet and exercise: - as monotherapy when	
metformin is considered inappropriate due to intolerance or contraindications - in addition to other medicinal products for the	Jun 2024
treatment of diabetes. SMC restriction: in addition to other oral anti-diabetic medicines as an option when glucagon-like peptide-1 (GLP-1) receptor agonists would be considered.	
SMC 2653	
Indication: for weight management, including weight loss and weight maintenance, as an adjunct to a reduced-calorie diet and increased physical activity in adults with an initial Body Mass Index (BMI) of ≥30 kg/m² (obesity) or ≥27 kg/m² to <30 kg/m² (overweight) in the presence of at least one weight-related comorbid condition (e.g., hypertension, dyslipidaemia, obstructive sleep apnoea, cardiovascular disease, prediabetes, or type 2 diabetes mellitus).	Jun 2024
SMC restriction: for use in adults with BMI ≥30 kg/m ^{2*} and at least one weight-related comorbidity.	
*a lower BMI cut-off may be more appropriate for members of minority ethnic groups known to be at equivalent risk of the consequences of obesity at a lower BMI than the white population.	
Xonvea® 10 mg/10 mg gastro-resistant tablets (doxylamine succinate/pyridoxine hydrochloride)	
FG1 450/22	
Indication: Severe nausea and vomiting in pregnancy (NVP) and Hyperemesis gravidarum (HG)	ТВС

END