

SMC Drug ID	Conditions	Decision	Date published on SMC Website	Date of decision / Expected date of decision
SMC2614	as monotherapy for the treatment of adult patients with relapsed or refractory (R/R) diffuse large B-cell lymphoma (DLBCL), after two or more lines of systemic therapy.	Available in line with local or regional guidance	10/06/2024	31/08/2024
Other Decision	n Specified :			

		Decision	Date published on SMC Website	Date of decision / Expected date of decision
	treatment of haemolytic anaemia due to sickle cell disease (SCD) in adults and paediatric patients 12 years of age and older as monotherapy or in combination with hydroxycarbamide. SMC restriction: as a second line treatment for haemolytic anaemia in patients with SCD who are intolerant, ineligible or have an inadequate response to, hydroxycarbamide.	Not routinely available as not recommended for use in NHS Scotland	10/06/2024	31/08/2024
Other Decision	Specified:			

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SMC Drug ID	Conditions	Decision	Date published on SMC Website	Date of decision / Expected date of decision
SMC2632	as monotherapy for the treatment of adult patients with relapsed or refractory (R/R) diffuse large B-cell lymphoma (DLBCL) after two or more lines of systemic therapy.	Available in line with local or regional guidance	10/06/2024	31/08/2024
Other Decision	n Specified :			

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	Date of decision / Expected date of decision	Date published on SMC Website	Decision	Conditions	SMC Drug ID
treated with ruxolitinib.	31/08/2024	10/06/2024		adult patients with moderate to severe anaemia who have primary myelofibrosis, post polycythaemia vera myelofibrosis or post essential thrombocythaemia myelofibrosis and who are	SMC2636
Other Decision Specified :				Specified:	Other Decision

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SMC Drug ID	Conditions	Decision	Date published on SMC Website	Date of decision / Expected date of decision
SMC2653	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). SMC restriction: for use in adults with BMI ≥30 kg/m2* 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.	Not routinely available as the ADTC is waiting for further advice from local clinical experts	10/06/2024	31/08/2024
Other Decision	n Specified :			

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SMC Drug ID	Conditions	Decision	Date published on SMC Website	Date of decision / Expected date of decision
SMC2655	for the treatment of patients 16 years of age and older with moderately to severely active ulcerative colitis (UC) who have had an inadequate response, lost response, or were intolerant to either conventional therapy, or a biological agent.	Not routinely available as there is a local preference for alternative medicines	10/06/2024	31/08/2024
Other Decisio	n Specified :			
Web Link: ht website.pdf	tps://www.scottishmedicines.org.uk/media/8383/etrasimod-film-c	coated-tablets-velsipity-abb-final-may-2024-for-		

clostridium botulinum neurotoxin type A (Xeomin®)					
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SMC2680	focal spasticity of the lower limb affecting the ankle joint.	Not routinely available as not recommended for use in NHS Scotland	10/06/2024	19/06/2024	
Other Decision					
Web Link: htt	ps://www.scottishmedicines.org.uk/media/8380/clostridium-botu	linum-neurotoxin-type-a-xeomin-non-sub-final-may-2024-			

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SMC Drug ID	Conditions	Decision	Date published on SMC Website	Date of decision / Expected date of decision
SMC2681	as monotherapy for the treatment of adult patients with newly diagnosed acute myeloid leukaemia (AML) who are ineligible for standard induction chemotherapy.	Not routinely available as not recommended for use in NHS Scotland	10/06/2024	19/06/2024
Other Decision	n Specified : Non-submission			

SMC Drug ID	Conditions	Decision	Date published on SMC Website	Date of decisio / Expected date of decision
SMC2682	treatment of eosinophilic esophagitis in adults and adolescents 12 years and older, weighing at least 40 kg, who are inadequately controlled by, are intolerant to, or who are not candidates for conventional medicinal therapy.	Not routinely available as not recommended for use in NHS Scotland	10/06/2024	19/06/2024
Other Decision	n Specified: Non-submission			
Web Link: ht	tps://www.scottishmedicines.org.uk/media/8382/dupilumab-dupix	ent-non-sub-final-may-2024-for-website.pdf		

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SMC Drug ID	Conditions	Decision	Date published on SMC Website	Date of decision / Expected date of decision
SMC2683	in combination with gemcitabine and cisplatin for the first-line treatment of locally advanced unresectable or metastatic biliary tract carcinoma in adults.	Not routinely available as not recommended for use in NHS Scotland	10/06/2024	19/06/2024
Other Decisio	n Specified: Non-submission			
Web Link: ht	tps://www.scottishmedicines.org.uk/media/8385/pembrolizumab-k	keytruda-non-sub-final-may-2024-for-website.pdf		

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