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 Decisio	n Specified :			

SMC2626 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. Available in line with national guidance 10/06/2024 31/08/2024 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. 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. Movies and paediate in the second s	SMC Drug ID	Conditions	Decision	Date published on SMC Website	Date of decision / Expected date of decision
have an inadequate response to, hydroxycarbamide.	SMC2626	(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	Available in line with national guidance	10/06/2024	31/08/2024
Other Decision Specified :		1			
	Other Decisio	n Specified :			

	b (Tepkinly)			
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.	Not routinely available as the ADTC is waiting for further advice from local clinical experts	10/06/2024	31/08/2024
Other Decisio	n Specified :			
Web Link : ht	tps://www.scottishmedicines.org.uk/media/8387/epcoritamab-tepl	kinly-final-may-2024-amended-050624-for-website.pdf		
momelotini	h (Omijara)			
	O an little and	Desite test	Determined Patrick	Defended and the
SMC Drug ID	Conditions	Decision	Date published on SMC Website	Date of decision / Expected date of decision
	Conditions 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.	Available in line with local or regional guidance		/ Expected date
SMC Drug ID	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.	Available in line with local or regional guidance	on SMC Website	/ Expected date of 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 \geq 30 kg/m2 (obesity) or \geq 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 \geq 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 Decisio	n Specified :			
Web Link • ht	tps://www.scottishmedicines.org.uk/media/8389/tirzepatide-moun	iaro-final-may-2024-amended-050624-for-website odf		

etrasimod (
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 the ADTC is waiting for further advice from local clinical experts	10/06/2024	31/08/2024
Other Decision	n Specified :			
Web Link : <u>ht</u>	tps://www.scottishmedicines.org.uk/media/8383/etrasimod-film-c	coated-tablets-velsipity-abb-final-may-2024-for-		
Web Link : <u>ht</u> website.pdf	•	coated-tablets-velsipity-abb-final-may-2024-for-		
Web Link : <u>ht</u> website.pdf	tps://www.scottishmedicines.org.uk/media/8383/etrasimod-film-c	oated-tablets-velsipity-abb-final-may-2024-for-	Date published on SMC Website	Date of decision / Expected date of decision
Web Link : <u>ht</u> website.pdf Clostridium	tps://www.scottishmedicines.org.uk/media/8383/etrasimod-film-constructions.org.uk/media/8383/etrasimod-film-constructi			/ Expected date
Web Link : <u>ht</u> website.pdf Clostridium SMC Drug ID SMC2680	tps://www.scottishmedicines.org.uk/media/8383/etrasimod-film-c botulinum neurotoxin type A (Xeomin®) Conditions	Decision Not routinely available as not recommended for use in	on SMC Website	/ Expected date of decision

uechabine (edazuridine (Inaqovi)			
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			
	ps://www.scottishmedicines.org.uk/media/8381/decitabine-cedaz	zuridine-inaqovi-non-sub-final-may-2024-for-website.pdf		
Web Link : <u>ht</u> dupilumab (SMC Drug ID	· · · · · · · · · · · · · · · · · · ·	Decision	Date published on SMC Website	Date of decision / Expected date of decision
dupilumab (Dupixent®)			/ Expected date
dupilumab (SMC Drug ID SMC2682	Dupixent®) Conditions 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	Decision Not routinely available as not recommended for use in	on SMC Website	/ Expected date of decision

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-ł	evtruda-non-sub-final-may-2024-for-website.pdf		