

SMC Drug ID	Conditions	Decision	Date published on SMC Website	Date of decision / Expected date of decision
SMC2641	For the treatment of adult and paediatric patients with vision loss due to inherited retinal dystrophy caused by confirmed biallelic RPE65 mutations and who have sufficient viable retinal cells.	Routinely available from a specialist centre in another health board	08/07/2024	30/09/2024
Other Decisio	n Specified :			
Web Link: ht	tps://www.scottishmedicines.org.uk/media/8464/voretigene-nepa	rvovec-luxturna-umar-final-june-2024-for-website.pdf		

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SMC Drug ID	Conditions	Decision	Date published on SMC Website	Date of decision / Expected date
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SMC2642	in adults for the treatment of chronic kidney disease.	Available in line with local or regional guidance	08/07/2024	30/09/2024
	SMC restriction: in patients having individually optimised			
	standard care (including angiotensin converting enzyme			
	inhibitors or angiotensin II receptor blockers, unless these are contraindicated or not tolerated), and either, at the start of			
	treatment:			
	an estimated glomerular filtration rate (eGFR) of 20			
	mL/min/1.73m2 up to 45 mL/min/1.73m2, or			
	an eGFR of 45 mL/min/1.73m2 up to 90 mL/min/1.73m2 and either:			
	o A urine albumin-to-creatinine ratio (uACR) of 22.6 mg/mmol			
	or more, or			
	o Type 2 Diabetes Mellitus (T2DM).			
Other Decisio	n Specified :			
Web Link · ht	tps://www.scottishmedicines.org.uk/media/8466/empagliflozin-jaro	diance-final-june-2024-amended-240624-for-website.ndf		

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SMC Drug ID	Conditions	Decision	Date published on SMC Website	Date of decision / Expected date of decision
SMC2644	in combination with trastuzumab, fluoropyrimidine and platinum-containing chemotherapy for the first-line treatment of locally advanced unresectable or metastatic HER2-positive gastric or gastro-oesophageal junction adenocarcinoma in adults whose tumours express PD-L1 with a CPS ≥ 1.	Not routinely available as not recommended for use in NHS Scotland	08/07/2024	19/06/2024
	n Specified : The submitting company's justification of the treat n addition the company did not present a sufficiently robust ecor			
	readment the company did not present a same lently result elec-	ionic analysis to gain acceptance by sivic.		
	tps://www.scottishmedicines.org.uk/media/8460/pembrolizumab-	, , , , ,		
Web Link: ht		, , , , ,		
Web Link: <a href="https://htt</td><td>tps://www.scottishmedicines.org.uk/media/8460/pembrolizumab-</td><td>, , , , ,</td><td>Date published on SMC Website</td><td>Date of decision / Expected date of decision</td></tr><tr><td>Meb Link: <a href=" htt<="" https:="" td=""><td>tps://www.scottishmedicines.org.uk/media/8460/pembrolizumab-relatlimab (Opdualag®)</td><td>keytruda-mgc-final-june-2024-for-website.pdf Decision Available in line with local or regional guidance</td><td>-</td><td>/ Expected date</td>	tps://www.scottishmedicines.org.uk/media/8460/pembrolizumab-relatlimab (Opdualag®)	keytruda-mgc-final-june-2024-for-website.pdf Decision Available in line with local or regional guidance	-	/ Expected date
Web Link: ht	relatlimab (Opdualag®) Conditions first-line treatment of advanced (unresectable or metastatic) melanoma in adults and adolescents 12 years of age and older.	keytruda-mgc-final-june-2024-for-website.pdf Decision Available in line with local or regional guidance	on SMC Website	/ Expected of decision

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pembrolizumab (Keytruda®)					
SMC Drug ID	Conditions	Decision	Date published on SMC Website	Date of decision / Expected date of decision	
SMC2660	in combination with fluoropyrimidine and platinum-containing chemotherapy, for the first-line treatment of locally advanced unresectable or metastatic human epidermal growth factor 2 (HER2)-negative gastric or gastro-oesophageal junction adenocarcinoma in adults whose tumours express programmed death-ligand 1 (PD-L1) with a combined positive score (CPS) ≥ 1.	Not routinely available as the ADTC is waiting for further advice from local clinical experts	08/07/2024	30/09/2024	
Other Decision	Specified:				
Web Link: htt	ps://www.scottishmedicines.org.uk/media/8461/pembrolizumab-k	keytruda-final-june-2024-for-website.pdf			

pegunigalsidase alfa (Elfabrio®)					
SMC Drug ID	Conditions	Decision	Date published on SMC Website	Date of decision / Expected date of decision	
SMC2665	for long-term enzyme replacement therapy in adult patients with a confirmed diagnosis of Fabry disease (deficiency of alpha-galactosidase). SMC restriction: for use in adults with symptomatic Fabry disease who would usually be offered an enzyme replacement therapy.	Available in line with national guidance	08/07/2024	08/07/2024	
Other Decision	n Specified :				
Web Link: htt	tps://www.scottishmedicines.org.uk/media/8459/pegunigalsidase	-alfa-elfabrio-final-june-2024-for-website.pdf			

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controlled ovarian stimulation for the development of mult follicles in women undergoing assisted reproductive technologies (ART) such as an in vitro fertilisation (IVF) or intracytoplasmic sperm injection (ICSI) cycle. SMC restriction: for use in normal responders (patients with	Not routinely available as there is a local preference for alternative medicines	08/07/2024	30/09/2024
anti-Müllerian hormone level of > 5.4 pmol/L) or high responders (patients with an anti-Müllerian hormone level ≥25 pmol/L).			
Other Decision Specified :			

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SMC Drug ID	Conditions	Decision	Date published on SMC Website	Date of decision / Expected date of decision
SMC2691	Film-coated tablets: in combination with other antiretroviral(s) for the treatment of adults with multidrug resistant HIV-1 infection for whom it is otherwise not possible to construct a suppressive anti-viral regimen, for oral loading prior to administration of long-acting lenacapavir injection. Solution for injection: in combination with other antiretroviral(s) for the treatment of adults with multidrug resistant HIV-1 infection for whom it is otherwise not possible to construct a suppressive anti-viral regimen.	Not routinely available as not recommended for use in NHS Scotland	08/07/2024	19/06/2024
Other Decision	n Specified: Non-submission			

SMC Drug ID	Conditions	Decision	Date published on SMC Website	Date of decision / Expected date of decision
SMC2692	in adults for intravenous induction and maintenance of general anaesthesia.	Not routinely available as not recommended for use in NHS Scotland	08/07/2024	19/06/2024
Other Decision	n Specified: Non-submission.			
Web Link: https://www.scottishmedicines.org.uk/media/8462/remimazolam-byfavo-non-sub-final-june-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
SMC2693	as monotherapy for the treatment of adult patients with advanced HER2-positive gastric or gastroesophageal junction (GEJ) adenocarcinoma who have received a prior trastuzumab-based regimen.	Not routinely available as not recommended for use in NHS Scotland	08/07/2024	19/06/2024
Other Decisio	n Specified: Non-submission			
Web Link: ht	tps://www.scottishmedicines.org.uk/media/8463/trastuzumab-der	uxtecan-enhertu-non-sub-final-iune-2024-for-website.pdf	:	

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