SMC Drug ID	Conditions	Decision	Date published on SMC Website	Date of decision / Expected date of decision
SMC2664	as monotherapy for the treatment of adult patients with locally advanced or metastatic cholangiocarcinoma with an isocitrate dehydrogenase-1 (IDH1) R132 mutation who were previously treated by at least one prior line of systemic therapy.	Not routinely available as the ADTC is waiting for further advice from local clinical experts	09/09/2024	15/11/2024
Other Decisio	n Specified :			
Web Link : ht	tps://scottishmedicines.org.uk/media/8577/ivosidenib-tibsovo-fina	I-august-2024-for-website.pdf		
dabrafenib	(Finlee®)			
SMC Drug ID	Conditions	Decision	Date published on SMC Website	Date of decision / Expected date of decision
SMC2667	in combination with trametinib (Spexotras®) for:	Not routinely available as the ADTC is waiting for further advice from local clinical experts	09/09/2024	15/11/2024
	 the treatment of paediatric patients aged 1 year and older with low-grade glioma with a BRAF V600E mutation who require systemic therapy. the treatment of paediatric patients aged 1 year and older with high-grade glioma with a BRAF V600E mutation who have received at least one prior radiation and / or chemotherapy treatment. 			
Other Decisio	 with low-grade glioma with a BRAF V600E mutation who require systemic therapy. the treatment of paediatric patients aged 1 year and older with high-grade glioma with a BRAF V600E mutation who have received at least one prior radiation and / or chemotherapy treatment. 			

	(Tecvayli)			
SMC Drug ID	Conditions	Decision	Date published on SMC Website	Date of decision / Expected date of decision
SMC2668	as monotherapy for the treatment of adult patients with relapsed and refractory multiple myeloma, who have received at least three prior therapies, including an immunomodulatory agent, a proteasome inhibitor, and an anti-CD38 antibody and have demonstrated disease progression on the last therapy.	Not routinely available as the ADTC is waiting for further advice from local clinical experts	10/09/2024	15/11/2024
Other Decisio	n Specified :			
	-			
Web Link : htt	ps://scottishmedicines.org.uk/media/8572/teclistamab-tecvavli-fit	nal-august-2024-for-website.pdf		
Web Link : <u>ht</u>	ps://scottishmedicines.org.uk/media/8572/teclistamab-tecvayli-fit	nal-august-2024-for-website.pdf		
Web Link : <u>ht</u>	ps://scottishmedicines.org.uk/media/8572/teclistamab-tecvayli-fi	nal-august-2024-for-website.pdf		
	ps://scottishmedicines.org.uk/media/8572/teclistamab-tecvayli-fin	nal-august-2024-for-website.pdf		
		nal-august-2024-for-website.pdf Decision	Date published on SMC Website	Date of decision / Expected date of decision
elranatamat	o (Elrexfio®)			/ Expected date
elranatamak SMC Drug ID SMC2669	(Elrexfio®) Conditions as monotherapy for the treatment of adult patients with relapsed and refractory multiple myeloma, who have received at least three prior therapies, including an immunomodulatory agent, a proteasome inhibitor, and an anti-CD38 antibody and have demonstrated disease progression on the last therapy.	Decision Not routinely available as the ADTC is waiting for	on SMC Website	/ Expected date of decision
elranatamak SMC Drug ID SMC2669 Other Decision	(Elrexfio®) Conditions as monotherapy for the treatment of adult patients with relapsed and refractory multiple myeloma, who have received at least three prior therapies, including an immunomodulatory agent, a proteasome inhibitor, and an anti-CD38 antibody and have demonstrated disease progression on the last therapy.	Decision Not routinely available as the ADTC is waiting for further advice from local clinical experts	on SMC Website	/ Expected date of decision

pegcetacop	lan (Aspaveli®)			
SMC Drug ID	Conditions	Decision	Date published on SMC Website	Date of decision / Expected date of decision
SMC2715	as monotherapy in the treatment of adult patients with paroxysmal nocturnal haemoglobinuria (PNH) who have haemolytic anaemia.	Not routinely available as not recommended for use in NHS Scotland	09/09/2024	21/08/2024
Other Decisio	n Specified : Non-submission			
	en (Waylivra®)	veli-non-sub-final-august-2024-for-website.pdf		
		veli-non-sub-final-august-2024-for-website.pdf Decision	Date published on SMC Website	Date of decision / Expected date of decision
volanesorse	en (Waylivra®)			/ Expected date
volanesorse SMC Drug ID SMC2716	en (Waylivra®) Conditions as an adjunct to diet in adult patients with genetically confirmed familial chylomicronaemia syndrome (FCS) and at high risk for pancreatitis, in whom response to diet and	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
SMC2717	as an add-on to standard therapy for the treatment of generalised myasthenia gravis (gMG) in adult patients who are anti-acetylcholine receptor (AChR) antibody positive.	Not routinely available as not recommended for use in NHS Scotland	09/09/2024	21/08/2024
Other Decisio	n Specified : Non-submission			
Web Link : ht	tps://scottishmedicines.org.uk/media/8574/zilucoplan-zilbrysg-no	n-sub-final-august-2024-for-website.pdf		