

fosdenopterin (Nulibry®)					
SMC Drug ID	Conditions	Decision	Date published on SMC Website	Date of decision / Expected date of decision	
SMC2624	Treatment of patients with molybdenum cofactor deficiency (MoCD) Type A.	Not routinely available as the ADTC is waiting for further advice from local clinical experts	13/01/2025	07/02/2025	
Other Decision	Specified: Ultra-orphan initial assessment				
Web Link: htt	ps://scottishmedicines.org.uk/media/8856/umar-fosdenopterin-r	nulibry-final-august-2024-issued-061224-for-website.pdf			

SMC Drug ID	Conditions	Decision	Date published on SMC Website	Date of decision / Expected date of decision
SMC2666	in adult women of reproductive age for symptomatic treatment of endometriosis in women with a history of previous medical or surgical treatment for their endometriosis.	Not routinely available as there is a local preference for alternative medicines	13/01/2025	20/08/2025
Other Decision Specified :				
Web Link: ht				

Page 1 of 6 27 August 2025 13:57:54



danicopan (Voydeya®)

NHS New Medicines Decisions for January 2025

SMC Drug ID	Conditions	Decision	Date published on SMC Website	Date of decision / Expected date of decision
SMC2675	as an add-on to ravulizumab or eculizumab for the treatment of adult patients with paroxysmal nocturnal haemoglobinuria (PNH) who have residual haemolytic anaemia.	Available in line with national guidance	13/01/2025	07/02/2025
	SMC restriction: under the advice of the national PNH service.			
Other Decision	n Specified :			
Web Link: ht	tps://scottishmedicines.org.uk/media/8849/danicopan-voydeya-fi	nal-dec-2024-for-website.pdf		
iptacopan (I SMC Drug ID	Fabhalta®) Conditions	Decision	Date published on SMC Website	Date of decision / Expected date of decision
	As monotherapy in the treatment of adult patients with paroxysmal nocturnal haemoglobinuria (PNH) who have haemolytic anaemia.	Decision Not routinely available as the ADTC is waiting for further advice from local clinical experts		/ Expected date
SMC Drug ID SMC2676	As monotherapy in the treatment of adult patients with paroxysmal nocturnal haemoglobinuria (PNH) who have haemolytic anaemia. SMC restriction: under the advice of the national PNH service.	Not routinely available as the ADTC is waiting for	on SMC Website	/ Expected date of decision
SMC Drug ID	As monotherapy in the treatment of adult patients with paroxysmal nocturnal haemoglobinuria (PNH) who have haemolytic anaemia. SMC restriction: under the advice of the national PNH service.	Not routinely available as the ADTC is waiting for	on SMC Website	/ Expected date of decision

Page 2 of 6 27 August 2025 13:57:54



risankizumab (Skyrizi®)
SMC Drug ID Conditions

NHS New Medicines Decisions for January 2025

				of decision
SMC2686	for the treatment of adult patients with moderately to severely active ulcerative colitis who have had an inadequate response to, lost response to, or were intolerant to conventional therapy or a biologic therapy.	Not routinely available as the ADTC is waiting for further advice from local clinical experts	13/01/2025	07/03/2025
Other Decision	n Specified :			
Web Link: htt	tps://scottishmedicines.org.uk/media/8852/risankizumab-skyrizi-a	abb-final-dec-2024-for-website.pdf		
		·		
sirolimus (H	lyftor®)			
sirolimus (H SMC Drug ID	dyftor®) Conditions	Decision	Date published on SMC Website	Date of decision / Expected date of decision
	· · ·	Decision Not routinely available as the ADTC is waiting for further advice from local clinical experts		/ Expected date
SMC Drug ID	For the treatment of adult and paediatric patients aged 6 and above with facial angiofibroma (facial AF) associated with tuberous sclerosis complex (TSC).	Not routinely available as the ADTC is waiting for	on SMC Website	/ Expected date of decision

Decision

Date published

on SMC Website / Expected date

Date of decision

Page 3 of 6 27 August 2025 13:57:54



		/ Expected date of decision
SMC2721 treatment of Duchenne muscular dystrophy (DMD) in patients aged 4 years and older.	le from a specialist centre in another 13/01/2025	07/03/2025
Other Decision Specified :		

SMC Drug ID	Conditions	Decision	Date published on SMC Website	Date of decision / Expected date of decision
SMC2728	as monotherapy for the treatment of adult and paediatric patients 12 years of age or older with a weight of 40 kg and above with paroxysmal nocturnal haemoglobinuria (PNH): In patients with haemolysis with clinical symptom(s) indicative of high disease activity. In patients who are clinically stable after having been treated with a complement component 5 (C5) inhibitor for at least the past 6 months. SMC restriction: under the advice of the national PNH service.	Not routinely available as the ADTC is waiting for further advice from local clinical experts	13/01/2025	07/03/2025
Other Decision	n Specified :			

Page 4 of 6 27 August 2025 13:57:54



ublituximab (Briumvi®)					
SMC Drug ID	Conditions	Decision	Date published on SMC Website	Date of decision / Expected date of decision	
SMC2731	treatment of adult patients with relapsing forms of multiple sclerosis (RMS) with active disease defined by clinical or imaging features.	Available in line with local or regional guidance	13/01/2025	07/03/2025	
	SMC restriction: treatment of relapsing-remitting multiple sclerosis (RRMS) with active disease defined by clinical or imaging features.				
Other Decision	Specified :				
Web Link: htt	ps://scottishmedicines.org.uk/media/8855/ublituximab-briumvi-a	abb-final-dec-2024-for-website.pdf			

ciclosporin (Cequa®)					
SMC Drug ID	Conditions	Decision	Date published on SMC Website	Date of decision / Expected date of decision	
SMC2739	treatment of moderate-to-severe Dry Eye Disease (keratoconjunctivitis sicca) in adult patients who have not responded adequately to artificial tears. SMC restriction: severe keratitis in adult patients with Dry Eye Disease.	Not routinely available as there is a local preference for alternative medicines	13/01/2025	07/03/2025	
Other Decision	Other Decision Specified :				
Web Link: htt	tps://scottishmedicines.org.uk/media/8847/ciclosporin-cequa-abb	p-final-dec-2024-for-website.pdf			

Page 5 of 6 27 August 2025 13:57:54



Web Link: https://scottishmedicines.org.uk/media/8853/rozanolixizumab-rystiggo-non-sub-final-dec-2024-for-website.pdf

bictegravir/e	emtricitabine/tenofovir alafenamide (Biktarvy®)			
SMC Drug ID	Conditions	Decision	Date published on SMC Website	Date of decision / Expected date of decision
SMC2760	Treatment of human immunodeficiency virus-1 (HIV-1) infection in paediatric patients at least 2 years of age and weighing at least 14 kg to less than 25kg without present or past evidence of viral resistance to the integrase inhibitor class, emtricitabine or tenofovir.	Not routinely available as not recommended for use in NHS Scotland	13/01/2025	18/12/2024
Other Decision	Specified: Non-submission			
2024-for-websi	ps://scottishmedicines.org.uk/media/8859/bictegravir-emtricitabir te.pdf Imab (Rystiggo®)	ne-tenofovir-alafenamide-biktarvy-non-sub-final-dec-		
		Decision	Data published	Date of decision
SMC Drug ID	Conditions	Decision	Date published on SMC Website	
SMC2761	as an add-on to standard therapy for the treatment of generalised myasthenia gravis (gMG) in adult patients who are anti-acetylcholine receptor (AChR) or anti-muscle-specific tyrosine kinase (MuSK) antibody positive.	Not routinely available as not recommended for use in NHS Scotland	13/01/2025	18/12/2024
Other Decision	1 Specified: Non-submission			

Page 6 of 6 27 August 2025 13:57:54