Enquiries to Ministry of Health 0800 855 066

## APPLICATION FOR SUBSIDY BY SPECIAL AUTHORITY

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APPLICANT (stamp or sticker acceptable)	PATIENT NHI:	REFERRER Reg No:			
Reg No:	First Names:	First Names:			
Name:	Surname:	Surname:			
Address:	DOB:	Address:			
	Address:				
Fax Number:		Fax Number:			
Antiretrovirals					
Initial application — Confirmed HIV Applications only from a named specialist. Approvals valid without further renewal unless notified.  Prerequisites(tick box where appropriate)  The patient has confirmed HIV infection Note: Tenofovir disoproxil prescribed under endorsement for HIV is included in the count of up to 4 subsidised antiretrovirals.  Subsidies apply for a combination of up to four antiretroviral medications. The combination of a protease inhibitor and low-dose ritonavir given as a booster (either as part of a combination product or separately) will be counted as one protease inhibitor for the purpose of accessing funding to antiretrovirals.					
Renewal — Confirmed HIV  Current approval Number (if known):					
Prerequisites (tick box where appropriate)	and value without latitude for lower arrivate from the				
The treatment remains appropriate and the patient is benefiting from treatment					
Initial application — Prevention of maternal transmission Applications only from a named specialist. Approvals valid for 1 year.  Prerequisites(tick boxes where appropriate)  Prevention of maternal foetal transmission or Treatment of the newborn for up to eight weeks					
Note: Tenofovir disoproxil prescribed under endorsement for HIV is included in the count of up to 4 subsidised antiretrovirals.  Subsidies apply for a combination of up to four antiretroviral medications. The combination of a protease inhibitor and low-dose ritonavir given as a booster (either as part of a combination product or separately) will be counted as one protease inhibitor for the purpose of accessing funding to antiretrovirals.  Some antiretrovirals are unapproved or contraindicated for this indication. Practitioners prescribing these medications should exercise their own skill, judgement, expertise and discretion, and make their own prescribing decisions with respect to the use of a Pharmaceutical for an indication for which it is not approved or contraindicated.					

I confirm the above details are correct and that in signing this form I understand I may be audited.

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Reg No:			First Names:	First Names:				
Name:				Surname:	Surname:			
Address:			DOB:	Address:				
				Address:				
Fax Nu	umbe	er:			Fax Number:			
Antir	etro	vira	ls - continued					
Initial application — post-exposure prophylaxis following exposure to HIV Applications from any relevant practitioner. Approvals valid for 4 weeks.  Prerequisites(tick boxes where appropriate)								
	and		Treatment course to be initiated wi	thin 72 hours post exposure				
				ad condomless anal intercourse or receptive vaginal intercourse with a known HIV positive person with an unknown viral load greater than 200 copies per ml				
		or						
		or	Patient has had non-consen required	sensual intercourse and the clinician considers that the risk assessment indicates prophylaxis is				
		or	Patient has had condomless is unknown	anal intercourse with a person from a high HIV previous	alence country or risk group whose HIV status			
a boo antire	ster trovi	(eithe rals.	er as part of a combination product	iretroviral medications. The combination of a proteas or separately) will be counted as one protease inhibit Society for HIV, Viral Hepatitis and Sexual Health Me				
			cond or subsequent post-exposu					
Applio	catio	ns fro	om any relevant practitioner. Approvitick boxes where appropriate)					
	and		Treatment course to be initiated wi	thin 72 hours post exposure				
			Patient has had condomless or detectable viral load great	s anal intercourse or receptive vaginal intercourse with ter than 200 copies per ml	h a known HIV positive person with an unknown			
		or	Patient has shared intravend	ous injecting equipment with a known HIV positive pe	rson			
		or	Patient has had non-consen required	sual intercourse and the clinician considers that the r	risk assessment indicates prophylaxis is			
		or	Patient has had condomless is unknown	anal intercourse with a person from a high HIV prev	alence country or risk group whose HIV status			
Applic	catio	ns on	ion — Percutaneous exposure lly from a named specialist. Approv tick box where appropriate)	rals valid for 6 weeks.				
Subsi	Ten dies ster	ofovi apply (eithe	y for a combination of up to four ant	to blood known to be HIV positive sement for HIV is included in the count of up to 4 sub irretroviral medications. The combination of a proteas or separately) will be counted as one protease inhibit	se inhibitor and low-dose ritonavir given as			

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Reg No:	First Names:	-				
Name:	Surname:	Surname:				
Address:	DOB:	Address:				
	Address:					
Fax Number:		Fax Number:				
Antiretrovirals - continued						
Renewal — Second or subsequent percutaneo	us exposure					
Current approval Number (if known):						
Applications only from a named specialist. Approvement of the properties of the prop	als valid for 6 weeks.					
The patient has percutaneous exposure to blood known to be HIV positive						

I confirm the above details are correct and that in signing this form I understand I may be audited.