## REPUBLIC OF SINGAPORE HEALTH SCIENCES AUTHORITY

**HEALTH PRODUCTS ACT 2007** 



## APPLICATION FOR CONSIGNMENT APPROVAL OF AN UNREGISTERED THERAPEUTIC PRODUCT FOR PATIENTS' USE

Please refer to the latest Guidance on HSA website before filling up the form. All applicants must comply with the Health Products Act (HPA) and its regulations.

SIGNED REQUEST FOR NAMED-	PATIENT APPLICATION TYPE (To b	pe completed by the requesting doctor or dentist)	
Purpose (Tick only one box)	☐ To import and supply an unregistered therapeutic product which presents a life-saving treatment option to a patient under my care whose condition will be clinicall compromised without the requested therapy, and that there is no effective alternative therapy registered in Singapore.		
	OR		
	☐ To import and supply a novel unregistered therapeutic product which offers a substantive clinical advantage over registered therapies and is expected to provide significant improvement in the clinical outcome of a patient under my care.		
	Note: Clinical justification(s) must be	provided.	
Conditions of Using this Special Access Route (Tick the box to indicate that you have read and agree to the conditions)	<ol> <li>The use of the unregistered therapeutic product is in compliance with Ministry of Health's allowable practice and applicable laws.</li> <li>The use of the unregistered therapeutic product is in compliance with the clinical practice allowed under the Singapore Medical or Dental Council's Ethical Code and Ethical Guidelines.</li> <li>Informed consent from the patient for the use of this unregistered therapeutic product will be obtained and documented.</li> </ol>		
Type of Application (Tick only one box)	□ New application	☐ Repeat application	
Number of Named-Patients (Please indicate a number)			
Product Name			
Dosage Form (Film-coated tablet, capsule, injection etc.)			
Strength (mcg, mg, mg/ml etc.)			
Required Quantity (Indicate quantity and unit of measure e.g. 3 boxes, 3 vials, 3 syringes etc.)			

Indicati	ion					
Dosage	e Regimen					
	n for Requesting for stered Therapeutic Product one box)	☐ The patient(s) have failed or tried registered therapies but there was inadequate response. Please list the registered therapies the patient(s) have failed or tried:  ☐ Other reasons, please state details:				
Supportive Evidence on the use of the Product in Named-Patient Applications  (Supportive evidence e.g. clinical practice guidelines or scientific literature should be submitted to support the use of the product, where appropriate.)		List the references submitted: 1. 2. 3. 4. 5.				
Particulars of Doctor or Dentist		Name:		Registration Number: (MCR or DCR number)		
		Department:				
		Practicing address:				
		Contact Number:		Email:		
		REQUESTER'S DECLARA	ATIONS	;		
	has not been evaluated for i I am fully aware that the co- unregistered therapeutic pro- I declare that I am fully respo I undertake to maintain reco- patient who received the uni I declare that all the informa- the information provided by i	fully aware that the therapeutic product requested in this application is not registered under the HPA and ot been evaluated for its quality, safety, and efficacy by the HSA.  fully aware that the consignment approval by HSA for my hospital/ clinic/ nursing home¹ to bring in the istered therapeutic product is not an endorsement of the clinical use by the Authority.  are that I am fully responsible for the use of the unregistered therapeutic product.  ertake to maintain records of the name, NRIC/identification document number and contact details of the nt who received the unregistered therapeutic product under my care.  are that all the information provided by me in this form is true and accurate. I acknowledge that if any of formation provided by me in this form is false or inaccurate, I will be liable to prosecution for providing false nation under the Penal Code.				
	Signature:			Date:		

Tick all boxes .

 $<sup>^{\</sup>rm 1}$  Specified healthcare service licensee under the Healthcare Services Act 2020.