

	MEDICAL DEVICES BRANCH REQUEST FOR UNREGISTERED MEDICAL DEVICE FOR USE ON PATIENTS BY QUALIFIED PRACTITIONER/ PHMC FACILITY
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* Please complete all fields below. All fields are mandatory.

Section A: Purpose of application

GN-26: On request by Qualified Practitioner for use on his patient	GN-27: On request by PHMC facility for use on their patients
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Section B: Requesting Qualified Practitioner (QP) information

To be completed by Head of Department (or equivalent) of PHMC for GN-27.

Full name		MCR or DCR Number	
Department		Designation	
Email		Tel no	
Name of Hospital/Clinic			
PHMC Licence No			
Address			

Section C: Clinical justification

Please select the appropriate clinical justification(s):

- Absence of alternative treatment option
- Available alternative treatments failed or deemed ineffective or unsuitable for patient according to professional judgement
- Unregistered medical device is needed to minimise disruption to the continued supply of a similar registered medical device
- Absence of registered alternatives or lack of a specific feature in registered medical device
- User's (doctor or dentist) familiarity or expertise
- Established medical device with history of safe use in a licensed private hospital or medical clinic

Please provide elaboration on the basis for the above selection:

Section D: Declaration

! **IMPORTANT**

1. I am fully aware that the medical device(s) specified in **attached SAR Device List** has not been evaluated by the Authority for the required quality, safety and efficacy standards for supply in Singapore.
2. The import and/or supply of the unregistered medical device(s) are required for the use of the patient(s) under my care/ patients of the PHMC facility and I undertake to assume full responsibility for such use and shall adhere to the conditions of approval.
3. I undertake to ensure the patient is appropriately informed prior to treatment and consents to the treatment.
4. I undertake to maintain records of the patient including the contact details of the patient who received the above medical device(s) under my care/ the care of the PHMC facility.
5. I will ensure that this medical device will be used or administered in accordance to its intended purpose and indications for use as stated in the product owner's instructions for use.
6. I undertake to indemnify the government against all actions, claims or proceedings in respect of any adverse event, injury to or death of any person whomsoever arising out of or in connection with the use of the above unregistered medical device.
7. I hereby declare that all the information provided by me in this form is true and accurate. I acknowledge that if any of the information provided by me in this form is false or inaccurate, I will be liable to prosecution for providing false information under the Penal Code.

Date _____ Signature of Qualified Practitioner/ Head of Department _____