Form No.: MDSA-HCP Effective Date: 1 April 2022



#### **MEDICAL DEVICES CLUSTER**

REQUEST FOR UNREGISTERED MEDICAL DEVICE FOR USE ON PATIENTS BY QUALIFIED PRACTITIONER/ LICENSED HEALTHCARE FACILITY

\* 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 licensed healthcare facility for use or their patients	n
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#### Section B: Requesting Qualified Practitioner (QP) information

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

Full name		MCR or DCR Number	
Department		Designation	
Email		Tel no	
Name of Hospital/Clinic			
PHMC/ HCSA Licence No			
Address			
Section C: Clinical justific	ation		
Please select the appropri	ate clinical justification(s):		
☐ Absence of alternative t	reatment option		
☐ Available alternative treatments failed or deemed ineffective or unsuitable for patient according to professional judgement			
☐ Unregistered medical de	evice is needed to minimise disruption to the contir	nued supply of a similar r	egistered medical device
☐ Absence of registered a	alternatives or lack of a specific feature in registere	d medical device	
☐ User's (doctor or dentist	t) familiarity or expertise		
☐ Established medical de	vice with history of safe use in a licensed private he	ospital or medical clinic	
Please provide elaboration	n 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 licensed healthcare facility and I undertake to assume full responsibility for such use.
- 3. I undertake to ensure the patient is appropriately informed prior to treatment and consents to the treatment.

Clinical Justification Review Form for unregistered Class D medical devices is included in this application

- 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 licensed healthcare 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	
		Page 1 of 2

Form No.: MDSA-HCP Effective Date: 1 April 2022



#### **MEDICAL DEVICES CLUSTER**

# REQUEST FOR UNREGISTERED MEDICAL DEVICE FOR USE ON PATIENTS BY QUALIFIED PRACTITIONER/ LICENSED HEALTHCARE FACILITY

Complete the below sections if the application is requested by a <u>public healthcare instituition (PHI)</u> and contain <u>Class C and/or Class D</u> medical devices.

# Section E: Endorsement by Chairman of Medical Board (CMB) or equivalent.

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

(!)	IMPORTANT
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I support the request of the unregistered Class C and/or Class D medical devices in this application

- 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 patients of the licensed healthcare facility.
- 3. 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 CMB or Equivalent