

## APPLICATION FOR APPROVAL TO PRESCRIBE MEDICINAL CANNABIS

Medicines Poisons and Therapeutic Goods Regulation 2008

## ALL DETAILS MUST BE COMPLETED AND ALL DETAILS MUST BE LEGIBLE OR CONSIDERATION OF THIS APPLICATION MAY BE DELAYED.

FAX to: 02 6205 0997 Please tick if application is URGENT  $\square$ **SECTION A: PRESCRIBER DETAILS** Name: AHPRA Speciality: **AHPRA Registration Number: Practice Name:** Phone: Fax: Email: **Practice Address: SECTION B: PATIENT DETAILS** Name: Date of Birth: \_\_\_\_/\_\_\_ Sex: ☐ M □F  $\square$  x Address: **DIAGNOSIS: INDICATION FOR CANNABIS: PROPOSED DOSE: EXPECTED MAXIMUM** DOSE: **ROUTE OF DELIVERY** ☐ VAPORISED ☐ ORAL LIQUID ☐ SPRAY ☐ OTHER (please specify)\_ IS A MEDICAL DEVICE TO □ мо **BE USED IN DELIVERY?:** YES (please specify) **SECTION C: APPROVAL BY DRUG SECTION C: CATEGORY APPROVAL** OR INDICATION: ☐ CATEGORY 6A (multiple sclerosis) ☐ CATEGORY 6B (nausea and vomiting related to cancer chemotherapy) **Complete Category** ☐ CATEGORY 6C (pain and/or anxiety in Approval or Approval patients with active malignancy or a life by Drug section only limiting disease where (in either case) the prognosis might reasonably be expected to be 12 months or less) ☐ CATEGORY 6D (refractory paediatric epilepsy)

COMPLETED FORMS TO BE SUBMITTED TO THE CHIEF HEALTH OFFICER

Phone: 02 6205 0998 Fax: 02 6205 0997 Email: <a href="mailto:hps@act.gov.au">hps@act.gov.au</a> Post: Locked Bag 5005, Weston Creek ACT 2611 #Please ensure all nosted applications are mark

#Please ensure all posted applications are marked confidential.



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SECTION D: PRODUCT	
NAME:	
MANUFACTURER NAME:	
MANUFACTURER ADDRESS:	
PRESENTATION:	□ VAPORISED
	☐ ORAL LIQUID
	□ SPRAY
	OTHER (please specify)
COMPOSITION	
(e.g. percentage composition of cannabinoids):	
SECTION E: SPECIALIST SUPPORT FOR USE	
NAME OF SPECIALIST:	
AHPRA SPECIALTY:	
SPECIALIST HAS:	☐ INITIATED CANNABIS THERAPY
	☐ PROVIDED WRITTEN OR VERBAL OPINION SUPPORTING CANNABIS THERAPY
SECTION F: ATTACHMENTS (please tick):	
☐ Product monograph or prescribing information from manufacturer.	
Authorised prescriber approval from the Therapeutic Goods Administration (TGA) (or confirmation that application has been submitted. Applications will not be approved by ACT Health until TGA approval has been obtained.	
PRESCRIBER DECLARATION (please tick):	
<ul> <li>I understand that the issue of an approval does not indicate Chief Health Officer support or endorsement of a proposed treatment.</li> <li>My patient is aware that this product has not been registered by the Therapeutic Goods Administration and should be considered experimental.</li> <li>My patient is aware that driving after taking medicinal cannabis may result in a positive drug driving test.</li> </ul>	
My patient is aware that driving after taking medicinal cannabis may result in a positive drug driving test.	
Signature of prescriber: Date:	
OFFICE USE ONLY	
Approved DAPIS Reference	e Number: Refused
Signature:	Date: Approval Term:/ to/

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