Australian Government

Department of Health

Office of Drug Control

Medicinal Cannabis Permit – Manufacture
MC0XX-S01-M01

This Medicinal Cannabis Permit is for the manufacture of a cannabis drug for one or more permitted supplies (as defined by the *Narcotic Drugs Act 1967* (**the Act**) and prescribed by the *Narcotic Drugs Regulation 2016* (the Regulation)) and is subject to the obligations set out in the Act and in the Regulation, and any conditions imposed by the Secretary on the Medicinal Cannabis Licence to which this Permit relates.

Licence holder:	MN Organisation Pty Ltd
Licence number:	MC0XX
Licensed premises:	1 Example Street Suburb STATE 0000
Specified area(s) of the licensed premises:	Room 123, Floor 1 Site A
Permit period:	28 January 2022 and 27 January 2023 inclusive

Authorised activities:

The licence holder is authorised, in accordance with this permit, to conduct activities at the specified area of the licensed premises to the following extent:

1. for one or more of the permitted supplies specified in the licence; and
2. in the quantities and within the limits as set out in Schedule 1: Manufacture Activities below.

Specified supply of a cannabis drug:

For the purposes of licence condition 2, the supply of a cannabis drug as identified in Column A of an item in the table in Schedule 2: Supply, in the circumstances described in Column B for that item, and which complies with the requirement (if any) in Column C for that item, is specified.

Signed: *Ann Example* Date: 28 January 2022

Ann Example

Delegate of the Secretary

Department of Health

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Instrument History

Version Number	Date	Description
Version 01	28 January 2022	Original grant

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Schedule 1: Manufacture Activities

			Quantity
Raw/Starting material	Cannabis	*Total quantity of cannabis authorised to be **obtained** over the life of this permit (kg) **(dry weight at 10% moisture content)**.*	1000
		*Maximum quantity of cannabis authorised onsite at **any one time (kg) (dry weight at 10% moisture content)**.*	150
	Cannabis extract	*Total quantity of cannabis extract authorised to be **obtained** over the life of this permit (kg).*	200
		*Maximum quantity of cannabis extract authorised onsite at **any one time (kg)**.*	50

			Quantity
Manufacture activities	Drugs manufactured from cannabis under this permit	*Total quantity of delta-9 tetrahyrdocannabinols (THC) **authorised to be manufactured** (kg).*	0
		*Maximum amount of delta-9 THC **authorised on the premises** at any one time (kg).*	0
		*Total quantity of non-delta-9 THC (THC Isomers other than delta-9 THC) **authorised to be manufactured** onsite (kg).*	0
		*Maximum amount of non-delta-9 THC (THC Isomers other than delta-9 THC) **authorised on the premises** at any one time (kg).*	0
		*Maximum amount of cannabidiol (CBD) **authorised to be manufactured** on site (kg).*	200
		*Maximum amount of CBD **authorised on the premises** at any one time (kg).*	50

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Schedule 2: Supply

ITEM	Column A SUPPLY	Column B CIRCUMSTANCES	Column C REQUIREMENT
1	Supply to another person (**recipient**)	The cannabis drug is registered goods for the purposes of the *Therapeutic Goods Act 1989*	The licence holder must make and keep a record of the ARTG number relevant to the registered goods
2	Supply to another person (**recipient**)	The recipient holds a medicinal cannabis licence granted under the *Narcotic Drugs Act 1967* which authorises the manufacture of a cannabis drug	The licence holder must make and keep a record of the name of the recipient
3	Supply to another person (**recipient)**	The supply of the cannabis drug is in accordance with an approval or authority under the *Therapeutic Goods Act 1989*	The licence holder must make and keep a record of the name of the recipient
4	Supply to another person (**recipient**)	The recipient holds a licence under Part 3-3 of the *Therapeutic Goods Act 1989*, and the cannabis drug is supplied for use by that person in the manufacture of a medicine (within the meaning of that Act)	The licence holder must make and keep a record of the name of the recipient
5	Supply by way of export	The licence holder has been issued an export licence and permission pursuant to the *Customs (Prohibited Exports) Regulations 1958*	The licence holder must make and keep a record of: • the name of the importing country; and • the export licence number

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ITEM	Column A SUPPLY	Column B CIRCUMSTANCES	Column C REQUIREMENT
6	Supply to another person (**recipient**)	The cannabis drug is supplied for use by a pharmacist in a public hospital for the purposes of that pharmacist dispensing the drug in accordance with the *Therapeutic Goods Act 1989*	The licence holder must make and keep a record of the name of the hospital and the responsible pharmacist
7	Supply to another person (**recipient**)	The cannabis drug is supplied for use as a reference standard for medical or scientific testing purposes	The licence holder must make and keep a record of the name of the recipient
8	Supply to another person (**recipient**)	The cannabis drug is supplied for use in a clinical trial that is, or is likely to be, approved (CTA) or notified (CTN) under the *Therapeutic Goods Act 1989*	The licence holder must make and keep a record of the CTA or CTN number
9	Supply to another person (**recipient**)	The cannabis drug is supplied for use in medical or scientific research which: (i) is not a clinical trial that is, or is likely to be, approved (CTA) or notified (CTN) under the *Therapeutic Goods Act 1989*; and (ii) does not involve the drug being administered to humans	The licence holder must obtain, and keep a record of, the details of the research and the manner in which the cannabis drug is to be used