



Australian Government
Department of Health
 Therapeutic Goods Administration

Public Summary

Summary for ARTG Entry: 292455 Microsafe Care Australia Pty Ltd - Nanocyn Disinfectant & Sanitiser with Specific Claims - Disinfectant, hospital grade

ARTG entry for Other Therapeutic Good - Registered disinfectant

Sponsor Microsafe Care Australia Pty Ltd
Postal Address 150 Williams Road, Mount Duneed, VIC, 3216
 Australia
ARTG Start Date 5/08/2017
Product Category Other Therapeutic Good
Status Active
Approval Area Medical Devices

Conditions

Conditions applicable to all therapeutic goods as specified in the document "Standard Conditions Applying to Registered or Listed Therapeutic Goods Under Section 28 of the Therapeutic Goods Act 1989" effective 1 July 1995.

Conditions applicable to the relevant category and class of therapeutic goods as specified in the document "Standard Conditions Applying to Registered or Listed Therapeutic Goods Under Section 28 of the Therapeutic Goods Act 1989" effective 1 July 1995.

Products

1 . Nanocyn Disinfectant & Sanitiser with Specific Claims - Disinfectant, hospital grade

Product Type	Single Device Product	Effective Date	9/08/2017 12:35:28 PM
GMDN	9950 Disinfectant, hospital grade		
Intended Purpose	Nanocyn Disinfectant & Sanitiser is a hard surface disinfectant effective against bacteria and viruses, including Staphylococcus aureus (MRSA), Herpes, Norovirus (Gastro) and Coronavirus, including SARS-CoV-2 (COVID-19), within 30 seconds. Not intended for use on medical devices or other therapeutic goods. Not to be used on skin.		

Specific Conditions

For the hard surface disinfectants that are retained on the part of the ARTG for registered goods, you, as sponsor, are required to comply with the standard and specific conditions for registered goods in Appendix 4, Standard Conditions applying to Registered or Listed Therapeutic Goods Under Section 28 of the Therapeutic Goods Act 1989 of DR4 applied at the time of registration and is ongoing.

AUST R

Pursuant to Section 28(2B) of the Therapeutic Goods Act 1989, I have decided to impose "The listing number must be placed on the label of the device by writing the number, immediately preceded by "AUST R" so that it is clearly visible to the user on (i) the label of the device; or (ii) the label on the outermost level of packaging in which the device is to be supplied to the user

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