VERIDIA

Certification Pack.

Kwiksan.

This pack contains
Product Information
TGA Certificate

Date Released

September 2020

Kwiksan

Hospital Grade Disinfectant
75% Ethanol sanitising spray
AUST L 342553

Kwiksan is a versatile, must-have, infection control solution for cleaning in environments where efficient and fast control against pathogenic micro-organisms is essential, and in areas where sensitivity to moisture is a critical factor. Kwiksan can be used on almost any surface including non-therapeutic devices, electronic equipment, trolleys, materials handling equipment and any other hard surface.

Ideal areas of application for Kwiksan are:

- Phones
- Keyboards, mouse, monitors,
- Headphones
- Touch screens and other electronic equipment.
- Printers and scanners
- Eftpos machines
- Glass, mirrors and shiny surfaces
- Push plates on doors
- Tools and equipment sensitive to water.
- And many more.

Easy to use
- Ready to use, non rinsing formula.
- Non-smear and non-streak benefit

Directions for use:
- Apply Kwiksan directly to surface.
- Polish dry with clean paper towel or a microfibre cloth.
- Kwiksan can also be spray applied to surfaces and left to air dry.

Safe
- Non-corrosive, non bleaching formula.
- Designed for safe use on electronic surfaces

Effective
- Rapid 60 second kill on COVID-19
- Contains a secondary biocide for a wider spectrum kill.
- Leaves a clean, streak-free surface that is thoroughly sanitised.
- Effective against gram negative and gram positive bacteria.
- Provides fast kill rate on non-therapeutic surfaces.
- Contains no quaternary ammonium compound.

Note
- Take extra care when disinfecting specialised electronic equipment. Pretest for chemical sensitivity and follow equipment manufacturers guidelines on cleaning / disinfecting methodology.

Fast germ killing in healthcare, aged care, veterinary and dental, child care, education and hospitality

Surfaces cleaned with Kwiksan are not only left sanitised and streak-free, but they will also stay clean for longer.

Actichem | 1800 108 800 | www.actichem.com.au
11 Gamma Close Beresfield NSW 2322 | info@actichem.com.au

Australian Government

Department of Health
Therapeutic Goods Administration

Australian Register of Therapeutic Goods Certificate

Issued to

Everard Paynter and Keith Paynter Family Trust

for approval to supply

Everard Paynter and Keith Paynter Family Trust - AP689 Kwiksan - Disinfectant, hospital grade

ARTG Identifier	342553
ARTG Start Date	28/08/2020
Product Category	Other Therapeutic Good Other Therapeutic Good - Listed disinfectant
Intended Purpose	Kwiksan is an alcohol based hospital grade disinfectant disinfectant for smooth surfaces and electronics. Effective against Pseudomanas. Aeruginosa, Proteus vulgaris, Escherichia coli,Staphylococcus aureus, salmonella choleraesuis and SARS-CoV-2 (COVID-19). Not to be used on skin. Not to be used on medical devices or other therapeutic goods.

Manufacturer Details	Address	Certificate number(s)
Applied Products Australia Pty Ltd	11 Gamma Close Beresfield , NSW , 23 22 Australia	

ARTG Standard Conditions

The above Other Therapeutic Good Other Therapeutic Good - Listed disinfectant has been entered on the Register subject to the following 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 Covered by This Entry

1. AP689 Kwiksan - Disinfectant, hospital grade

Product Specific Conditions

- 1. Standards
  The listed goods must comply with standards applicable to those goods under part 3 of the Act.
- 2. Changes to Goods
  Changes to Goods Changes or variations in respect of any information concerning the listed therapeutic goods, being information that would have been relevant* to a decision to list the goods in the Register, including information on the formulation of the listed goods or other aspects of their manufacture, and the labelling of the goods, shall forthwith be notified to the Secretary, or the Secretary's delegate appointed for the purposes of section 28 of the Act, and where necessary*, the change or variation shall not be implemented until approved by the Secretary. (*Reference should also be made to the Guidance on the Regulation of Disinfectants in Australia).
- 3. Records Held
  i. The sponsor of the listed goods shall keep such records relating to the goods as are necessary: (a) to expedite recall if necessary of any batch of the listed goods; (b) to identify the manufacturer(s) of each batch of the listed goods. Where any part of or step in the manufacture in Australia of the listed goods is sub-contracted to a third party who is not the sponsor, copies of relevant Good Manufacturing Practice agreements relating to such manufacture shall be kept.
  ii. Each sponsor shall retain records of the distribution of all of the sponsor's listed goods for a period of five years and upon the request of the Therapeutic Goods Administration, shall provide the records or copies of the records.

- 4. Sampling
  The sponsor of the listed goods shall permit officers who have been authorised under the Regulations to do so to take samples of therapeutic goods and carry out related duties in accordance with the Regulations.
- 5. Overseas Regulatory Actions
  Where the listed goods are distributed regularly overseas as well as in Australia, product recall or any actions other similar regulatory action taken in relation to the goods outside Australia which has or may have relevance to the quality, safety or efficacy of the goods distributed in Australia must be notified to the Therapeutic Goods Administration via Post Market Surveillance email, MedicalDeviceSurveillance@health.gov.au as soon as the action or information is known to the sponsor.
- 6. Indications
  In relation to listed goods, the sponsor must have and shall retain, while the goods remain listed, evidence necessary to substantiate and support the accuracy of the indications in relation to the listed goods and, upon the request of the Therapeutic Goods Administration, shall produce such evidence.

Therapeutic Goods Administration
PO Box 100, Woden ACT 2606 Australia
Phone: 1800 020 653
Email: info@tga.gov.au

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ARTG Start Date: 28/08/2020