

Department of Health

Therapeutic Goods Administration

Australian Register of Therapeutic Goods Certificate

Issued to

Ebos Group Australia Pty Ltd

for approval to supply

Ebos Group Australia Pty Ltd - Oximeter, pulse

ARTG Identifier 321974

ARTG Start date 19/08/2019

Product Category Medical Device Included Class IIa

GMDN 17148

GMDN Term Oximeter, pulse

Intended Purpose The Fingertip Pulse Oximeter, for the non invasive measurement of

SpO2 oxygen saturation and blood pulse rate.

Manufacturer Details	Address	Certificate number(s)
Shenzhen Aeon Technology Co Ltd	RM6H02 Block 27-29 Tianxia IC Industrial Park Majialong No 133 of Yiyuan Road Nantou Street, Nanshan District Shenzhen, 518052 China	DV-2019-MC-10702-1

ARTG Standard Conditions

The above Medical Device Included Class IIa has been entered on the Register subject to the following conditions:

- The inclusion of the kind of device in the ARTG is subject to compliance with all conditions placed or imposed on the ARTG entry. Refer Part 4-5, Division 2 (Conditions) of the Therapeutic Goods Act 1989 and Part 5, Division 5.2 (Conditions) of the Therapeutic Goods (Medical Devices) Regulations 2002 for relevant information.
 - Breaching conditions of the inclusion related to the device of the kind may lead to suspension or cancellation of the ARTG entry; may be a criminal offence; and civil penalties may apply.

Products Covered by This Entry

1. Oximeter, pulse

Product Specific Conditions

No specific conditions have been recorded against this entry.

Therapeutic Goods Administration PO Box 100, Woden ACT 2606 Australia

Phone: 1800 020 653 Email: info@tga.gov.au ARTG Identifier: 321974 ARTG Start Date: 19/08/2019