



Australian Government
Department of Health, Disability and Ageing
Therapeutic Goods Administration

Australian Register of Therapeutic Goods Certificate

Issued to

Philips Electronics Australia Ltd

for approval to supply

Philips Electronics Australia Ltd - Transducer, toco

ARTG Identifier	490453
ARTG Start Date	30/05/2025
Product Category	Medical Device Included Class IIb
GMDN	37258
GMDN Term	Transducer, toco
Intended Purpose	The device is intended to be used for monitoring and recording of, and to generate alarms for multiple physiological parameters of pregnant women and the fetus.

Manufacturer Details	Address	Certificate number(s)
Philips Medizin systeme Boblingen GmbH	Hewlett-Packard Strasse 2 BOBLINGEN , Boblingen , 71034 Germany	DV-2021-MC-18949-1

ARTG Standard Conditions

The above Medical Device Included Class IIb 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. Transducer, toco

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: 490453
ARTG Start Date: 30/05/2025



Australian Government
Department of Health, Disability and Ageing
Therapeutic Goods Administration

Australian Register of Therapeutic Goods Certificate

Issued to

Philips Electronics Australia Ltd

for approval to supply

**Philips Electronics Australia Ltd - Patient monitor module,
multifunction**

ARTG Identifier	449246
ARTG Start Date	21/05/2024
Product Category	Medical Device Included Class IIb
GMDN	36553
GMDN Term	Patient monitor module, multifunction
Intended Purpose	Measures uterine activity externally and measures additionally the maternal pulse (non-invasive). It allows additional measurements: DECG, MECG or IUP.

Manufacturer Details	Address	Certificate number(s)
Philips Medizin systeme Boblingen GmbH	Hewlett-Packard Strasse 2 BOBLINGEN , Boblingen , 71034 Germany	DV-2024-MC-07603-1

ARTG Standard Conditions

The above Medical Device Included Class IIb 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. Patient monitor module, multifunction

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: 449246
ARTG Start Date: 21/05/2024



Australian Government
Department of Health, Disability and Ageing
Therapeutic Goods Administration

Australian Register of Therapeutic Goods Certificate

Issued to

Philips Electronics Australia Ltd

for approval to supply

**Philips Electronics Australia Ltd - Patient monitor module,
multifunction**

ARTG Identifier 449245
ARTG Start Date 21/05/2024
Product Category Medical Device Included Class IIb
GMDN 36553
GMDN Term Patient monitor module, multifunction
Intended Purpose Measures DECG, MECG or IUP.

Manufacturer Details	Address	Certificate number(s)
Philips Medizin systeme Boblingen GmbH	Hewlett-Packard Strasse 2 BOBLINGEN , Boblingen , 71034 Germany	DV-2024-MC-07603-1

ARTG Standard Conditions

The above Medical Device Included Class IIb 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. Patient monitor module, multifunction

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: 449245
ARTG Start Date: 21/05/2024