# Anesthesia Gas Monitoring IntelliVue G1 & IntelliVue G5



#### **Features**

- Continuous, simultaneous, real-time monitoring of respiratory and anesthetic gases.
- IntelliVue G1 is fully integrated with the Philips IntelliVue MP20/30/40/50 families of anesthesia patient monitors.
- IntelliVue G5 is fully integrated with the Philips IntelliVue MP40/50/60/70/80/90 families of anesthesia patient monitors.<sup>1</sup>
- Measurement of oxygen (optional for IntelliVue G1; standard for IntelliVue G5), carbon dioxide, nitrous oxide, halothane, isoflurane, enflurane, sevoflurane, desflurane.
- Manual anesthetic agent selection (IntelliVue G1).

- Automatic Agent Identification (IntelliVue G5) of up to two anesthetic agents.
- Inspiration and expiration values for all gases.
- Airway Respiration Rate (awRR), derived from CO<sub>2</sub> waveform.
- Minimum alveolar concentration values (MAC).
- Long-term stability offering easy and infrequent calibration.
- Individual alarm limits for each parameter.
- Quick access to the most commonly performed functions.

 $1. In \ the \ USA \ \& \ Canada, \ Intelli Vue \ G5$  is also integrated with MP 20/30.

**PHILIPS** 

# Description

The M1013A IntelliVue G1 and the M1019A IntelliVue G5 provide a non-dispersive infrared measurement of respiratory and anesthetic gases and a paramagnetic measurement of oxygen (Fast O<sub>2</sub>).

The IntelliVue G1 is designed to work with the Philips IntelliVue MP20/30/40/50 Anesthesia option #H30 and the IntelliVue G5 with the MP40/50/60/70/80/90<sup>1</sup> Anesthesia option #H30 through a digital interface (RS232).

Both gas analyzers are intended for measuring the airway gases of ventilated patients during the induction of, maintenance of, and emergence from anesthesia.

The gas analyzers produce display waves for  $O_2$ ,  $CO_2$ ,  $N_2O$ , and anesthetic agents, together with numerics for inspired and end-tidal values for  $O_2$ ,  $CO_2$ ,  $N_2O$ , anesthetic agents, MAC values and airway respiration rate.

With the IntelliVue G1 an anesthetic agent must be selected manually for measurement.

The IntelliVue G5 automatically identifies two out of five anesthetic agents at a time.

An automatic zero calibration is performed by the IntelliVue G1 and IntelliVue G5 as required to maintain measurement accuracy.

# **Specifications**

# **Physical**

#### Size (H x W x D)

85 x 300 x 232 mm, (3.35 x 11.81 x 9.13 inches).

#### Weight

< 4 kg (8.8 lb.) net.

#### **Safety**

Complies with:

- IEC/EN 60601-1
- UL 60601-1
- CSA C22.2 No. 601.1-M90
- IEC/EN 60601-1-1
- IEC/EN 60601-1-2

#### 1.In the USA and Canada, IntelliVue G5 is also integrated with MP20/30

#### Caution:

U.S. Federal Law restricts this device to sale by or on the order of a physician.

# **Environmental**

#### **Operating Temperature**

 $10 \text{ to } 40 \,^{\circ}\text{C}$  (50 to  $104 \,^{\circ}\text{F}$ ).

#### Storage

-20 to 65 °C (-4 to 149 °F).

#### **Operating Humidity**

5 to 90 % RH max at @ 40 °C (104 °F).

#### **Storage Humidity**

5 to 95 % RH max. at @ 65 °C (149 °F).

# **Operating Altitude**

-305 to 2900 m, (-1,000 to 9,515 ft.).

#### Storage Altitude

-305 to 5000 m, (-1,000 to 16,404 ft.).

# Sample Flow Rate

200 ml/minute (± 20 ml/min.).

# **Warmup Time**

1-2 minutes to measure CO<sub>2</sub>, typically less than 6 minutes for full accuracy specifications.

#### **Interface**

RS-232-C (RJ45)

#### **Calibration Interval**

Annual Gas calibration test. Calibration is stable and normally only requires checking.

# Units

mmHg, kPa, vol% for all gases,

(except for CO<sub>2</sub>, no vol%).

# **Power Requirements**

- Input: 100 to 240 V ±10 % ac.
- Frequency: 50/60 Hz.
- Power consumption: 45 VA max.

#### **Monitored Gases / Parameters**

- O<sub>2</sub>
- CO<sub>2</sub>
- Airway Respiration Rate (awRR)
- N<sub>2</sub>O
- The anesthetic agents Halothane, Enflurane, Isoflurane, Sevoflurane and Desflurane. IntelliVue G1 can measure one of these manually selected agents at a time. IntelliVue G5 can automatically detect two of these agents at a time.
- Minimum alveolar concentration (MAC) values

#### Oxygen Measurement

#### Range

5 to 100 vol%.

#### Accuracy<sup>1</sup>

± 3 vol%.

#### Resolution

1 vol%.

#### Rise-time

< 500 msec typical.

#### **Carbon Dioxide Measurement**

#### Range

0 to 76 mmHg.

#### Accuracy<sup>1</sup>

± 0.5 vol% or 12 % relative, which ever is greater.

#### Resolution

1 mmHg.

#### Rise-time

# 350 msec typical<sup>2</sup>

#### **Airway Respiration Rate**

Derived from CO<sub>2</sub> waveform.

#### Range

0 - 60 rpm.

#### Accuracy

± 1 rpm.

#### Resolution

1 rpm.

#### **Detection Criteria**

adaptive threshold

#### **Nitrous Oxide Measurement**

#### Range

0 to 100 vol%.

# Accuracy 1

2.0 % + 8 % relative.

#### Resolution

1 vol%.

#### Rise-time

500 msec typical <sup>2</sup>

# **Anesthetic Agent Measurement**

The IntelliVue G1 and IntelliVue G5 identify any of the following anesthetic agents.

For IntelliVue G1 the agent must be selected manually. IntelliVue G5 automatically identifies up to two of the following agents:

Table 1: Anesthetic Agent Measurement

Agent	Range (vol%)	Accuracy	Resolution (vol%)	Rise time typical (msec)
Halothane	0 - 8.5	0.15 vol% + 15.0 % relative	0.05	< 500
Enflurane	0 - 10.0	0.15 vol% + 15.0 % relative	0.05	< 500
Isoflurane	0 - 8.5	0.15 vol% + 15.0 % relative	0.05	< 500
Sevoflurane	0 - 10.0	0.15 vol% + 15.0 % relative	0.05	< 500
Desflurane	0 - 20.0	0.15 vol% + 15.0 % relative	0.05	< 500

<sup>1.</sup>Accuracy specifications refer to BTPS for mmHg kPa and STPD for vol% at 40-60 % relative humidity. They refer to M1658A tubing and M1657B watertrap.

<sup>2.</sup>With 2.6m M1658A tubing and M1657B watertrap (10-90 % step response) at 25°C and 40% relative humidity.

# **Sample Delay Time**

All measurements are subject to a delay of 5 seconds due to the length of the sampling path<sup>1</sup>.

# **Total System Response Time**

The Total System Response Time is the sum of Sample Delay Time and Rise Time.

#### **Zero Calibration**

The IntelliVue G1 and G5 request the first zero calibration not later than 5 minutes after power-on. In a steady state, a normal zero is performed not more than every 2 hours. The normal zero calibration lasts less than 21 seconds.

The  $\rm O_2$  zero calibration is performed once as part of the startup and typically once every 4 months of continuous operation / standby state. An  $\rm O_2$  zero calibration can last up to 60 seconds.

#### **Alarms**

#### **Limit Alarms**

Limit alarms are provided for the numeric values displayed by the IntelliVue G1 and IntelliVue G5 (see table 2 below).

# response) at 25 °C and 40 % Table 2: Alarm Limits

Gas	High Limit Range	Low Limit Range
etCO <sub>2</sub>	20 - 76 mmHg	10 - 75 mmHg
imCO <sub>2</sub>	0 - 20 mmHg	none
inN <sub>2</sub> O	0 - 82 vol%	none
inO <sub>2</sub> (Optional)	19 - 100 vol%	18 - 99 vol%
in/et HAL/ISO/ENF	0.1 - 7.5 vol%	0.0 - 7.4 vol%
in/et SEV	0.1 - 9.0 vol%	0.0 - 8.9 vol%
in/et DES	0.2 - 20.0 vol%	0.0 - 19.8 vol%
awRR	10 - 60 rpm (Adult, Pediatric) 30 - 60 rpm (Neonatal)	0 - 55 rpm

#### **Alarm Delay**

< 19 seconds if no automatic zero calibration occurs within that time.

#### **Apnea Alarm**

#### Adjustable Delay Range

10 - 40 seconds

#### **Alarm Criterion**

No detected breath within the adjusted delay time.

#### Apnea Alarm Delay

Within 2 seconds after this criterion is met, if no automatic zero calibration occurs.

#### **INOP Alarms**

Unique INOP messages and with or without an INOP tone are triggered on the host monitor if certain conditions occur.

The exact messages depend on the host monitor used and are described in the monitor's accompanying documentation.

<sup>1.</sup>With 2.6 m M1658A tubing and M1657B watertrap (10-90 % step response) at 25 °C and 40 % relative humidity.

# **Options**

The following options are available for the

#### IntelliVue G1

- #A01 without O<sub>2</sub> measurement
- #A02 with paramagnetic O<sub>2</sub> measurement

One of the following K-options must be ordered:

- #K11 IntelliVue Interface Cable (1.5 m)
- #K12 IntelliVue Interface Cable (3.0 m)

#### IntelliVue G5

One of the following K-options must be ordered:

- #K11 IntelliVue Interface Cable (1.5 m)
- #K12 IntelliVue Interface Cable (3.0 m)

#### Accessories

To ensure the specified performance of and to prevent possible damage to the M1013A IntelliVue G1 and the M1019A IntelliVue G5, **only** the following accessories may be used:

- M1658A Gas Sample Tube, 2.6 m
- M1657B Watertrap.
- 13902A Elbow Airway Adapter, 15 mm (right angle).
- M1612A Straight Airway Adapter.
- M1655A/B Gas Exhaust Return Line.<sup>1</sup>
- M1656A/B Gas Exhaust Return Filter. 1

For service purposes, the following accessories may be used:

- M1662A Calibration Gas
- M1659A Calibration Bag.

*Note:* These accessories are not included with the IntelliVue G1 or the IntelliVue G5.

#### **Documentation**

The User's Guide and Service and Installation Guide for this module are included in the documentation package shipped with the Philips IntelliVue Systems.

<sup>1.</sup> May not be available in all countries.

# Philips Medical Systems is part of Royal Philips Electronics

#### Interested?

Would you like to know more about our imaginative products? Please do not hesitate to contact us. We would be glad to hear from you.

On the web www.medical.philips.com

Via e-mail
medical@philips.com

*By fax* +31 40 27 64 887

By postal service
Philips Medical Systems
Global Information Center
P.O Box 1168
5602 BD Eindhoven
The Netherlands



The IntelliVue G1 (M1013A) and the IntelliVue G5 (M1019A) comply with the requirements of the Council Directive 93/42/EEC of 14 June 1993 (Medical Device Directive).

Asia

Tel: +852 2821 5888

Europe, Middle East, Africa Tel: +31 40 27 63005

Latin America
Tel: +55 11 2125 0764

*North America* Tel: +1 800 229 6417

© Koninklijke Philips Electronics N.V. 2004 All rights are reserved. Reproduction in whole or in part is prohibited without the prior written consent of the copyright holder.

Philips Medical Systems Nederland B.V. reserves the right to make changes in specifications or to discontinue any product at any time without notice or obligation and will not be liable for any consequences resulting from the use of this publication.

Printed in The Netherlands. 4522 962 11301 FEB 2006

