

Evita 2 dura



**Intensive Care Ventilator
Instructions for Use
Software 4.n**

Working with these Instructions for Use

Header line – the title... of the main chapter

The title of the specific sub-section is printed underneath the main header – to help you find your way quickly from subject to subject.

Page body... the Instructions for Use

in combined text/illustrations. The information is expressed in the form of practical actions, giving the user direct hands-on experience in learning how to use the machine.

Left-hand column... the text

provides explanations and instructs the user step-by-step in the practical use of the product, with short, clear instructions in easy-to-follow sequence.

Bullet points indicate separate actions. Where several actions are described, numbers are used both to refer to the relevant details in the illustrations and to specify the sequence of actions.

Right-hand column... the illustrations

provide the visual reference for the text and make it easier to locate the various parts of the equipment. Elements mentioned in the text are highlighted. Unnecessary details are avoided. Screen displays prompt the user to proceed and confirm correct actions.

Preparing for use
Fitting components

Preparing for use

The following instructions cover:

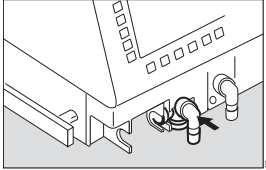
- Assembly of the various equipment items.
- Electrical connection and gas connection.
- Setting the language for display texts.
- Automatic device check with calibration of the sensors.

Fitting components

- Always use properly prepared parts, see "Care", page 90.

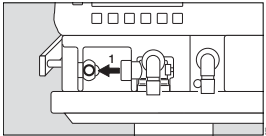
Fitting the expiration valve

- Push the expiration valve block firmly into the mounting. Check that it is properly secured by gently pulling the port.



Fitting the flow sensor

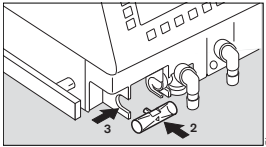
- 1 Push the socket to the left as far as it will go.



- 2 Insert the flow sensor – with the probe facing towards the ventilator – into the mounting and push it into the socket as far as it will go.

Then:

- 3 Push flow sensor to the right as far as it will go into the rubber lip of the expiration valve.



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What's new in Evita 2 dura software 4.n

Specification of the humidifier used

- »Active humidifier«
or
- »HME/Filter« (artificial nose)
- for more accurate measurement of the volume parameters

Apnoea ventilation On/Off

- can be selected as starting configuration

Extended range of settings for the alarm time T_{Apnoea} ✓^x

- from 5 to 60 seconds
(formerly 15 to 60 seconds)

Ventilation mode BIPAP_{Assist}

- for pressure-controlled assisted ventilation

Patient mode »prev. patient« can be selected

- to adopt the settings, including alarms, which were effective before switching off the equipment

Leakage compensation On/Off

- for activation and deactivation of the automatic leakage compensation function

Monitoring of tube blockages

- New alarm message »Tube blocked !!!«

Additional weaning parameters

available as software version 4.n plus upgrade
in addition to the parameter occlusion pressure P 0.1
Evita 2 dura 4.n also determines the parameters

- RSB Rapid Shallow Breathing index
and
- NIF Negative Inspiratory Force Index
- f_{spn} and MV_{spn} as trend

External flow source

available as software version 4.n plus upgrade

- The amount of external flow is calculated by Evita 2 dura 4.n (e.g. for additional tracheal gas insufflation) and adjusts the volume monitoring tolerances in order to avoid inadvertent alarms

Evita Remote (Remote Pad)

optionally available

- Remote control pad for parallel remote operation of function keys on Evita 2 dura 4.n

NIV

optionally available

- Application mode to support non-invasive ventilation therapies

Nurse call

optionally available

- Socket for connecting alarm signals to a central alarm station in the hospital

Simplified settings

under »other modes«

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For Your Safety and that of Your Patients

Strictly follow the Instructions for Use

Any use of the apparatus requires full understanding and strict observation of these instructions. The apparatus is only to be used for purposes specified here.

Maintenance

The apparatus must be inspected and serviced regularly by trained service personnel at six monthly intervals.

Repair and general overhaul of the apparatus may only be carried out by trained service personnel.

We recommend that a service contract be obtained with DrägerService and that all repairs also be carried out by them.

Only authentic Dräger spare parts may be used for maintenance.

Observe chapter "Maintenance Intervals".

Accessories

Do not use accessory parts other than those in the order list.

Note: Even reusable accessories (e.g. parts which can be cleaned and/or sterilized) have a limited useful life. Wear and tear may be increased and the service life reduced considerably as a result of various factors when handling and conditioning such parts (e.g. disinfectant residues left behind after autoclaving may corrode the material). Such parts must be replaced immediately if external signs of wear become evident, such as cracks, deformation, discoloration, peeling, etc.

Not for use in areas of explosion hazard

This apparatus is neither approved nor certified for use in areas where combustible or explosive gas mixtures are likely to occur.

Safe connection with other electrical equipment

Electrical connections to equipment which is not listed in these Instructions for Use should only be made following consultations with the respective manufacturers or an expert.

Liability for proper function or damage

The liability for the proper function of the apparatus is irrevocably transferred to the owner or operator to the extent that the apparatus is serviced or repaired by personnel not employed or authorised by DrägerService or if the apparatus is used in a manner not conforming to its intended use.

Dräger cannot be held responsible for damage caused by non-compliance with the recommendations given above.

The warranty and liability provisions of the terms of sale and delivery of Dräger are likewise not modified by the recommendations given above.

Dräger Medical GmbH

Safe use of the equipment

This equipment must only be used under the supervision of qualified medical staff, so that help is available immediately if any faults or malfunctions occur.

Medications containing alcohol and other flammable substances, must NOT be used in the patient circuit. This is a fire hazard!

Medicaments and other substances based on inflammable solvents, such as alcohol, must not be used in the patient system. Danger of fire!

Adequate ventilation must be ensured when using inflammable substances for disinfection.

Do not use mobile telephones within 10 metres of ventilators!

Mobile telephones may impair the functioning of electromedical equipment and endanger the patient *

When the unit is mounted onto the trolley it must NOT be tilted at more than 5°, this could cause the entire assembly to tip over.

Appropriate ventilation monitoring

The (built-in) monitoring facilities of the Evita 2 dura ventilator ensure appropriate monitoring of ventilation therapy and (therefore) will detect any undesirable changes in the following ventilation parameters:

- Airway pressure, P_{aw}
- Expiratory minute volume, MV
- Inspiratory O₂ concentration, FiO_2
- Inspiratory breathing gas temperature, T
- Expiratory CO₂ concentration, $etCO_2$ (optional)
- Inspiratory breathing volume, V_{Ti}
- Apnoea time
- Tachypnoea monitoring

Changes in these parameters may be caused by:

- Acute changes in the patient's condition
- Incorrect settings and faulty handling
- Equipment malfunctions
- Failure of power and gas supplies

If a fault occurs in this equipment, separate measuring instruments should be used.

Back-up ventilation with an independent manual ventilation device

If a fault is detected in Evita 2 dura so that its life-support functions are no longer assured, ventilation using an independent ventilation device must be started without delay – if necessary with PEEP and/or increased inspiratory O₂ concentration (e.g. with the Dräger Resutator 2000).

* Dräger medical equipment meets the requirements for immunity to interference in accordance with the specific product standards and EN 60601-1-2 (IEC 60601-1-2). Depending on the type of mobile telephone used and on the application situation, however, field strengths exceeding the values specified in the applicable standards may develop in the immediate vicinity of the mobile telephone and therefore lead to faults and malfunctions.

Intended Medical Application

Evita 2 dura

Long-term ventilator for intensive care.
For adults, children and neonates.

With the following ventilation modes

IPPV (Intermittent Positive Pressure Ventilation)
controlled and assisted constant-volume ventilation.

With the options:

- **CPPV (Continuous Positive Pressure Ventilation)**
- **PLV (Pressure Limited Ventilation)**
- **AutoFlow[®]** (optional)
for automatic regulation of inspiration flow
- **IRV (Inversed Ratio Ventilation)**

SIMV (Synchronized Intermittent Mandatory Ventilation)

Procedure for weaning patients off the ventilator after they have started spontaneous breathing.

With the options:

- **PLV (Pressure Limited Ventilation)**
- **AutoFlow[®]** (optional)
for automatic regulation of inspiration flow

MMV (Mandatory Minute Volume Ventilation)

Spontaneous breathing with automatic adjustment of mandatory ventilation to the patient's minute volume requirement.

With the options:

- **PLV (Pressure Limited Ventilation)**
- **AutoFlow[®]** (optional)
for automatic regulation of inspiration flow

SB (Spontaneous Breathing)

Spontaneous breathing at ambient pressure

CPAP (Continuous Positive Airway Pressure)

Spontaneous breathing with positive airway pressure

ASB (Assisted Spontaneous Breathing)

Pressure-assisted spontaneous breathing

BIPAP* (Biphasic Positive Airway Pressure)

Pressure-controlled ventilation combined with free spontaneous breathing during the complete breathing cycle, and adjustable pressure increase to CPAP level.

BIPAP^{Assist} (Biphasic Positive Airway Pressure Assisted)

Pressure-controlled assisted ventilation

APRV (Airway Pressure Release Ventilation) (optional)

Spontaneous breathing on two pressure levels with long time ranges – independently adjustable.

Special modes:

Apnoea Ventilation

For switching over automatically to volume-controlled mandatory ventilation, if breathing stops.

If apnoea occurs, Evita 2 dura emits an alarm after the preset alarm period ($T_{Apnoea} \sqrt{A}$) and starts volume-controlled ventilation.

ILV (optional)

Independent Lung Ventilation,

Separate, differentiated, synchronised ventilation with two units, one for each lung, either two Evita 2 dura units or an Evita 2 dura with another Evita unit.

Diagnostics

Intrinsic PEEP-measurement (optional)

for determining intrinsic PEEP and measuring trapped volume.

Occlusion pressure measurement (optional)

for evaluating breathing drive during spontaneous breathing.

With monitoring for

Airway pressure, P_{aw}

Expiratory minute volume, MV

Inspiratory O₂ concentration, FiO₂

Inspiratory breathing gas temperature, T

Expiratory CO₂ concentration, etCO₂ (optional)

Functional O₂ saturation and heart rate (optional)

Inspiratory breathing volume, V_{Ti}

Apnoea time


Tachypnoea monitoring to detect rapid, shallow spontaneous breathing

Automatic gas switch-over

In the event of a gas failure, the change-over to another gas is automatic.

* Registered trade mark

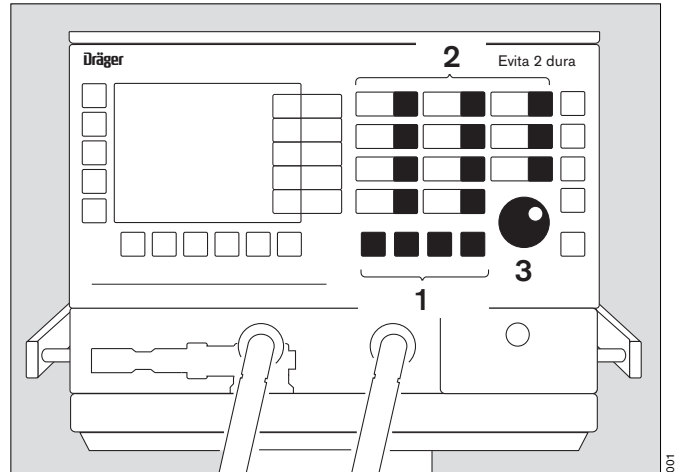
Operating Concept

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Operating Concept

Ventilation Controls

- 1 Keys for selecting the ventilation modes:
 - IPPV
 - SIMV
 - BIPAPand
 - **other modes**
- 2 Keys for selecting/setting ventilation parameters:
 - Tidal volume **V_T**
 - Inspiration time **T_{insp}**
 - Frequency **f**
 - Inspiratory flow **Flow**
 - Inspiratory pressure **P_{insp}**
 - Pressure-assisted spontaneous breathing **ΔPASB**
 - Positive end-expiratory pressure **PEEP**
 - Pressure rise time **Ramp**
 - O₂ concentration **O₂**
 - Sensitivity **Trigger**
- 3 Central "turn-and-push" rotary knob for setting the parameters:
 - To set = turn the rotary knob
 - To confirm setting = press the rotary knob.



Setting ventilation parameters









- 2 To set a ventilation parameter = press the corresponding parameter key. The yellow LED in the key lights up.
- 3 To set the value of the ventilation parameter = turn the rotary knob. The value is displayed next to the parameter key.
- 3 To confirm the value = press the rotary knob. The yellow LED goes out.

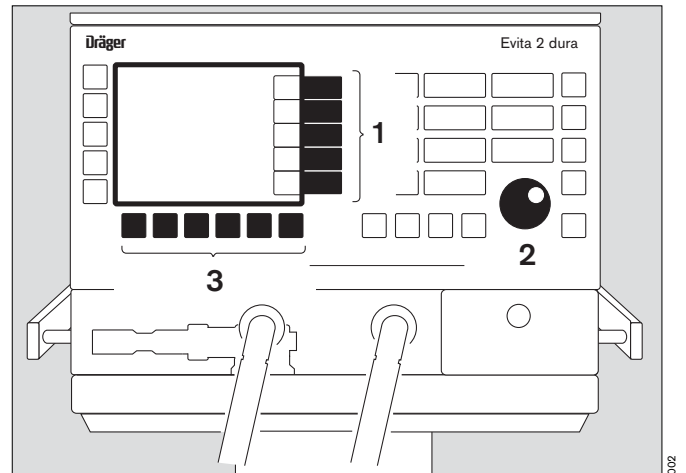
Selecting the ventilation mode

- 1 Hold down the appropriate key for about 3 seconds
or
press the appropriate key briefly and confirm = press the rotary knob.
The selected ventilation mode will now be activated.





For detailed instructions on setting the ventilation modes, see page 41.

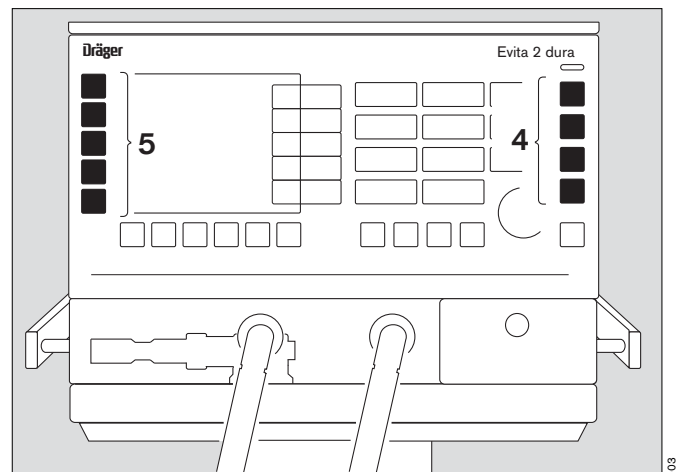
Screen Operating Controls

- 1 Menu keys for selecting the menu on the screen.
- 2 Central "turn-and-push" rotary knob for selecting and setting the options displayed on the screen.
To select/set = turn the rotary knob
To confirm = press the rotary knob.
- 3 Screen operating keys:
 - »Print  « key for manual printer logging,
 - »  /  « key for setting the screen brightness to bright or dark,
 - »Freeze  « key for freezing the curves,
 - »Curves   « key for displaying a different pair of curves,
 - »Values   « key for displaying a different combination of measured values,
 - one reserve key for future functions.



Keys for routine and additional functions

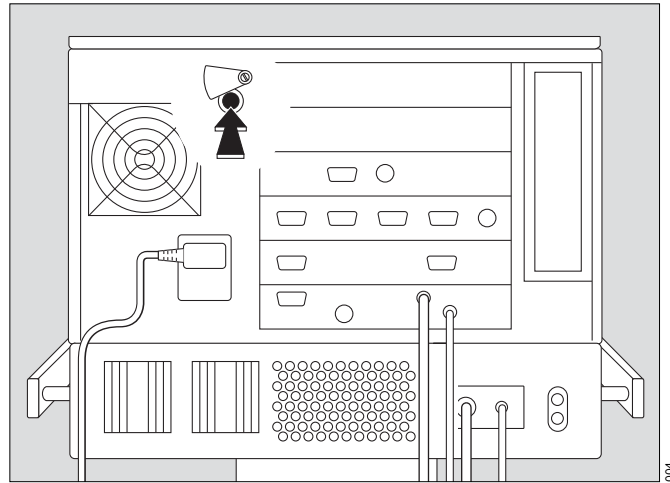
- 4 Frequently used keys for routine functions are positioned on the right-hand side of the front panel:
 - »  « key for suppressing the audible alarm tone,
 - »Alarm Reset« key for resetting or acknowledging alarm messages,
 - »  « key for calling up information and help on the desired setting,
 - »  « key for protecting against inadvertent or unauthorised modification of the settings of the ventilation parameters or ventilation modes.
- 5 Keys for additional functions are positioned on the left-hand side of the front panel:
 - »  « key for switching the medicament nebuliser on/off,
 - »O₂ ↑ Suction« key for oxygen enrichment during bronchial suction,
 - »Insp. hold« for manually activated inspiration,
 - »Exp. hold« for extending the expiration time,
 - one reserve key for future functions.



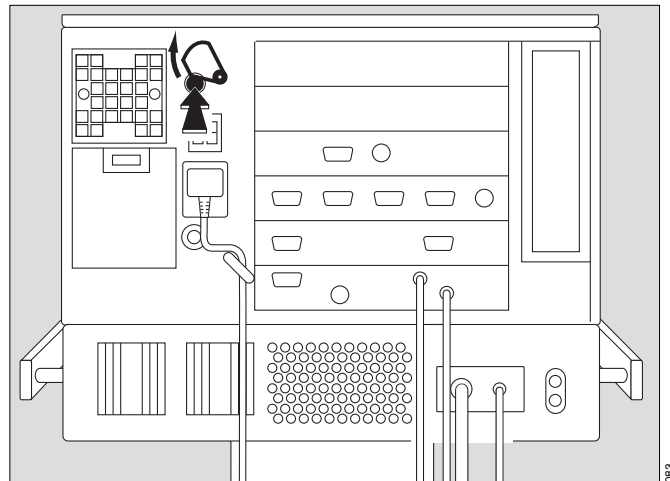
Power switch

To switch the apparatus on/off.

Located on the back panel, with a covering flap to protect the switch from being switched off inadvertently.



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Standby key »⏻«

Placed on its own, away from other keys.

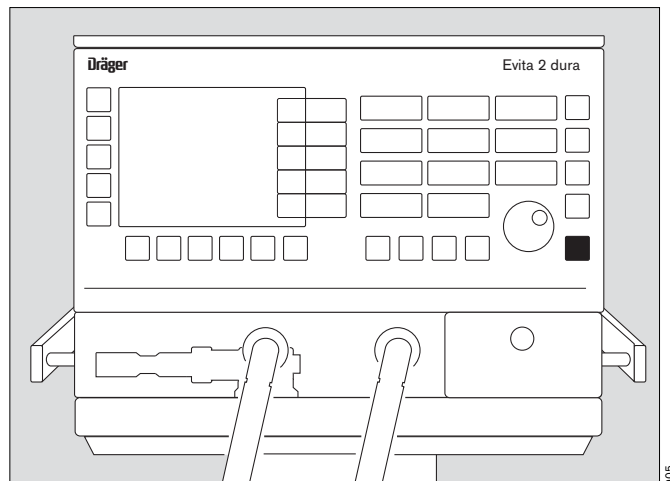
For keeping the apparatus on standby
or
for switching on ventilation.

To switch to standby:

- Press and hold down the »⏻« key for at least 3 seconds.

To switch on ventilation:

- Briefly press and release the »⏻« key.



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Screen Pages

The screen pages consist of two basic structures:

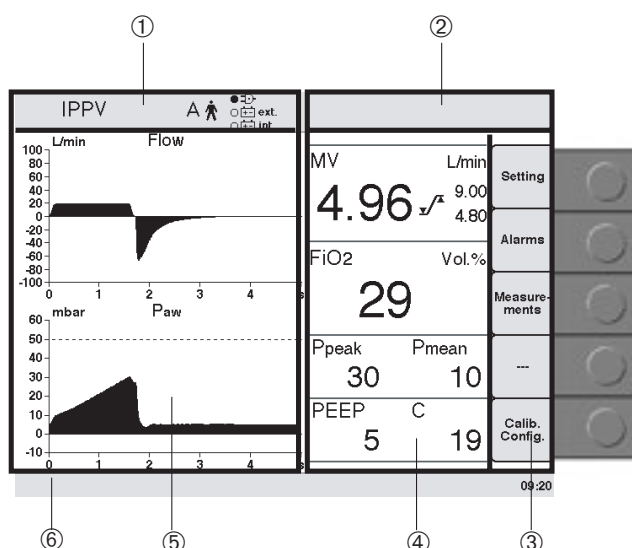
- the main page, displaying all important ventilation characteristics at a glance and
- application-specific pages for functions and settings.

Important functions are displayed in the same position in both structures:

- active ventilation mode and patient mode
- alarm, warning and advisory messages
- field for the menu selection keys
- information and help

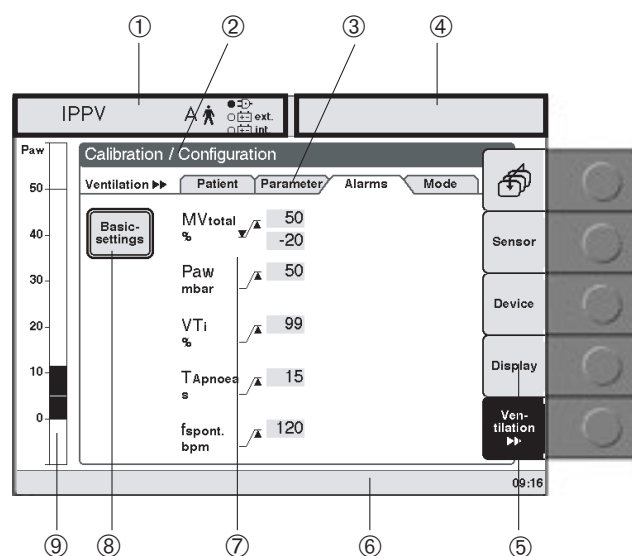
Structure of the main page

- ① Line for the active ventilation mode, patient mode and trigger indicator
- ② Bar for alarm, warning and advisory messages
- ③ Field for menu selection keys
- ④ Field for measured values
- ⑤ Field for curves
- ⑥ Bar for information and help



Structure of the application-specific pages

- ① Bar for the currently active ventilation mode and patient mode (example: BIPAP)
- ② Field for displaying the selected menu
- ③ Field for the menu bar
- ④ Bar for the alarm, warning and advisory messages
- ⑤ Field for menu selection keys
- ⑥ Bar for information and help
- ⑦ Screen field, selectable with the rotary knob
- ⑧ Screen key, selectable with the rotary knob
- ⑨ Field for continuous pressure display and monitoring

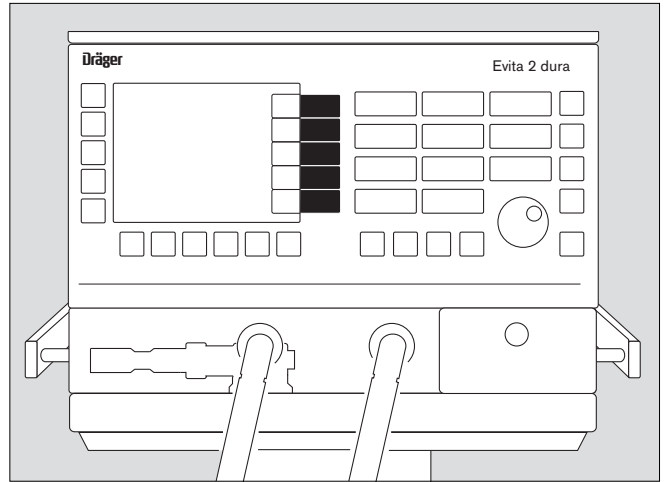


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The menu keys on the right-hand edge of the screen select the screen pages for the following specific application situations:

- Settings
- Alarms
- Measurements
- Measurement manoeuvre (optional)
- Calib./Config.



Screen page »Settings«

- For setting apnoea ventilation
- For setting intermittent PEEP (sigh)

For detailed operating instructions, see "Setting ventilation modes" on page 41 onwards.

Screen page »Alarms«

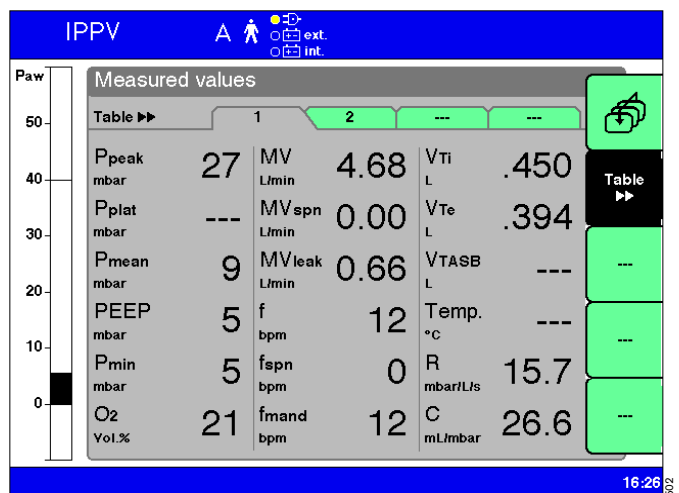
- For displaying the measured values with their alarm limits
- For setting the alarm limits

For detailed operating instructions, see "Setting alarm limits" on page 53.

Screen page »Measured values«

- For displaying all the measured values in the current ventilation mode.

Press the »Table ►►« screen key to display more option measured values in »Table 2«.



Screen page »Calibration/Configuration«

Sensors

- Calibrating the sensors for O₂ and flow
- Switching the monitoring system on and off

Device

- Setting the volume of the acoustic alarm
- Setting the screen contrast
- Setting the date and time
- Selecting language and measurement units
- Setting the external interfaces

Display

- Selecting 2 x 6 measured values from the main page
- Selecting 2 x 2 curves from the main page

Ventilation

- Patient mode
- Ventilation mode
- Ventilation parameters
- Alarm limits

Colour screen

For differentiating various items of information on the screen.

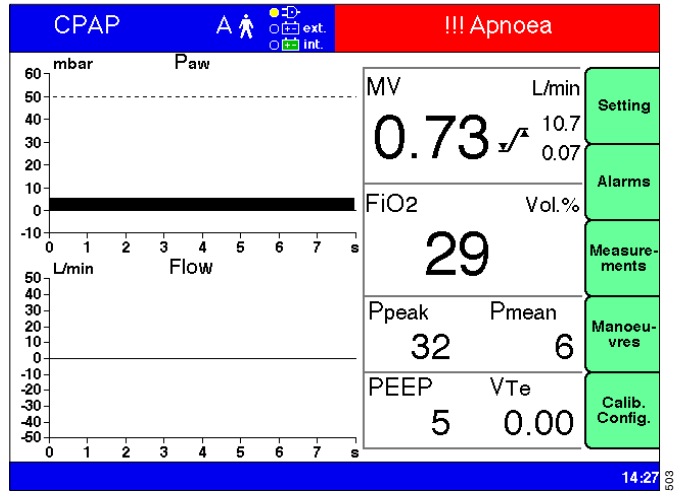
For messages

Red = Alarm

Yellow = Caution or advisory message

Blue = Alarm is no longer active

Example: »!!! Apnoea« alarm

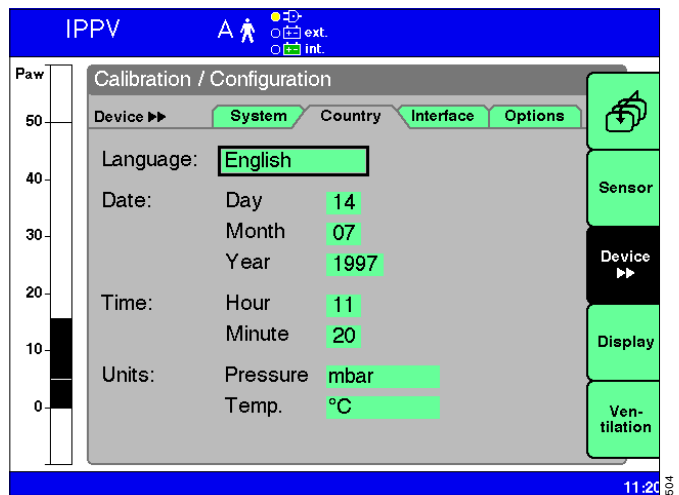


For menu buttons:

Green = Can be selected

Black = Has been selected

Example: Menu button »Device ▶▶«

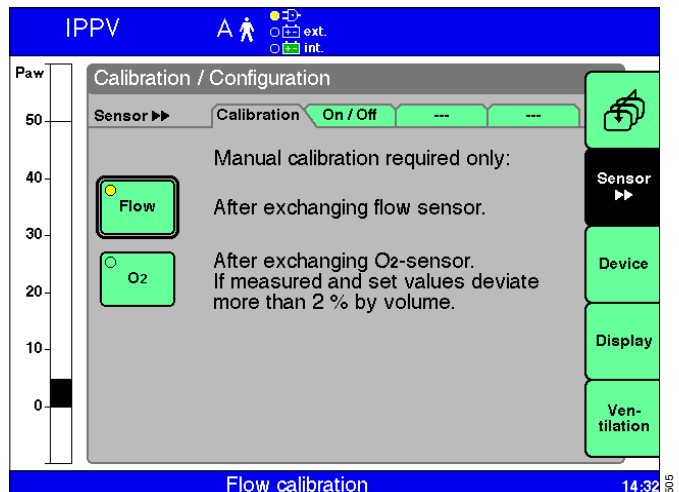


For screen keys

Green "LED" in the screen key = function not active

Yellow "LED" in the screen key = function active

Example: Screen key »Flow« = function active



Preparing

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Preparing for use

The following instructions cover:

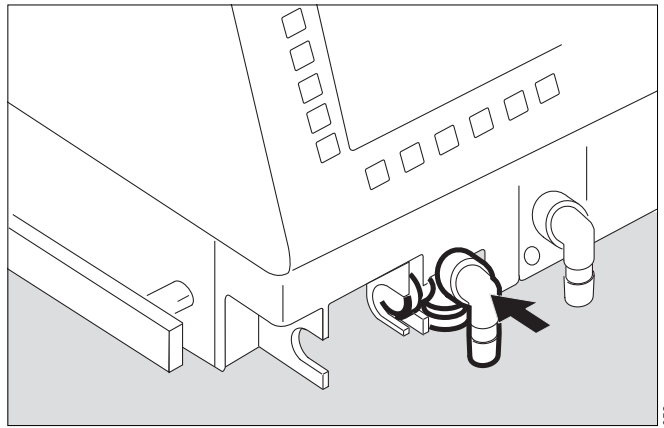
- Assembly of the various equipment items.
- Electrical connection and gas connection.
- Setting the language for display texts.
- Automatic device check with calibration of the sensors.

Fitting components

- Always use properly prepared parts, see "Care", page 90.

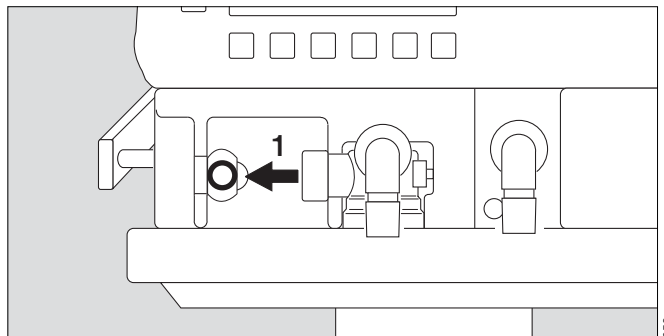
Fitting the expiration valve

- Push the expiration valve block firmly into the mounting. Check that it is properly secured by gently pulling the port.



Fitting the flow sensor

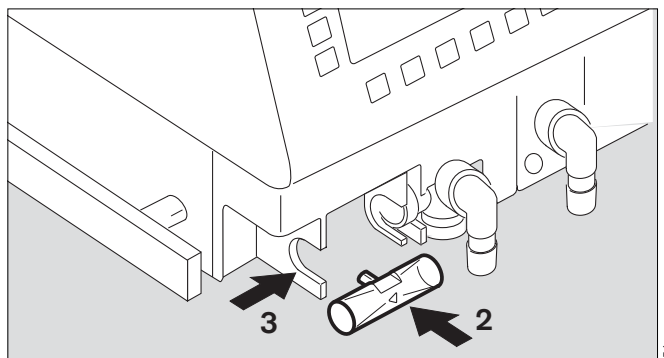
- 1 Push the socket to the left as far as it will go.



- 2 Insert the flow sensor – with the probe facing towards the ventilator – into the mounting and push it into the socket as far as it will go.

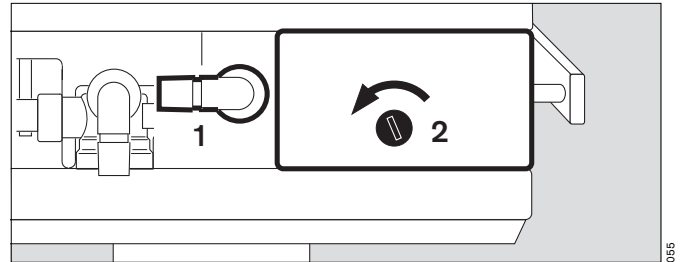
Then:

- 3 Push flow sensor to the right as far as it will go into the rubber lip of the expiration valve.

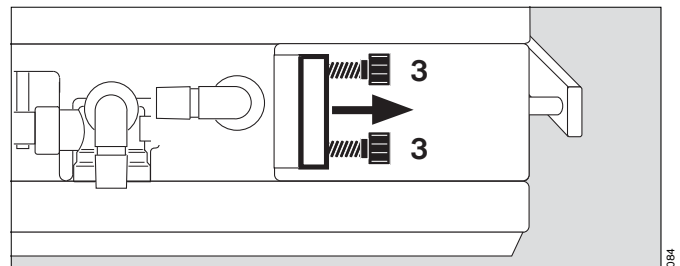


Fitting the O₂ sensor

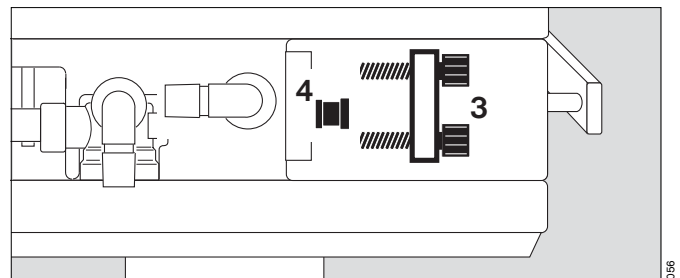
- when using the system for the first time
- when the display reads:
»O₂ measurement inop.«
- when calibration can no longer be performed.
- Tilt the control unit upwards.
- 1 Turn port downwards or to the left.
- 2 Unscrew (e.g. with coin), and remove protective cover.



- 3 Unscrew the two knurled screws and open the sensor housing.



- 4 Insert new sensor. The sensor end with the visible circular tracks on the contacts goes into the housing.
- 3 Close the sensor housing securely with the two knurled screws.
- Screw the protective cover back in place.



After inserting:

- Wait 15 minutes for the O₂ sensor to warm up. The O₂ sensor cannot be calibrated until it has warmed up.
- Calibrate O₂ sensor, page 66.
- Dispose of used sensor, page 99.

Note on the use of heat and moisture exchangers

The use of heat and moisture exchangers (HME's) in the patient connection can considerably increase breathing resistance. An increase in breathing resistance will in turn lead to greater effort in spontaneous breathing and/or greater trigger effort during assisted ventilation. Under unfavourable conditions, an increase in breathing resistance can lead to an unwanted intrinsic PEEP.

This breathing resistance in the patient connection cannot be directly monitored by the ventilator.

- The condition of the patient and the ventilator's measured values for air volume and resistance must be checked more frequently.
- Follow the Instructions for Use of the heat and moisture exchanger (HME).
- Do not use the heat and moisture exchanger (HME) at the same time as a medicament nebuliser or humidifier!

Note on the use of bacterial filters

The use of expiratory bacterial filters and/or HME filters on the ventilator is not recommended.

However, if bacterial filters are nevertheless used on the expiration side, an undesirable increase in breathing resistance is possible. Especially during medicament nebulisation and humidifying, the resistance of the expiratory bacterial filter may increase gradually. For the patient, the effect may be increased breathing effort and intrinsic PEEP.

An intrinsic PEEP can be recognised by the fact that the expiratory flow does not return to "0" before the end of expiration.

If PEEP is unacceptably high, the unit signals the »!!! PEEP high« alarm.

- Check the bacterial filter and replace it if it is the cause of the PEEP.

The inspiratory and expiratory breathing resistance of the patient system can be determined before ventilation by the device check in standby mode – see page 30 onwards.

For ventilating adults and children

From 100 mL tidal volume VT and above:

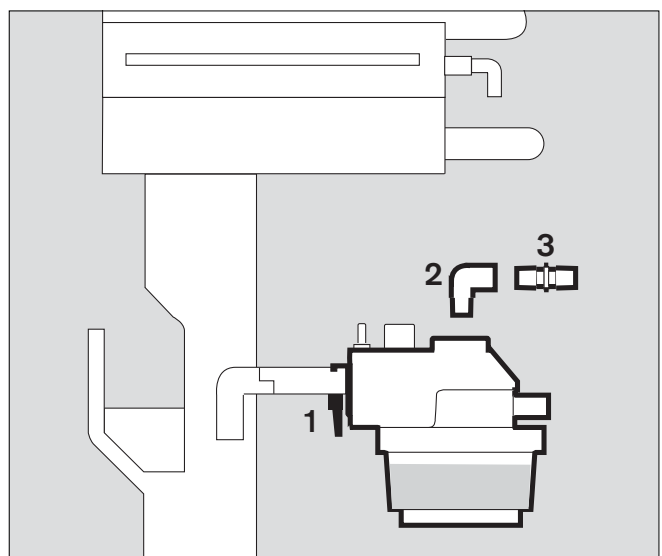
Patient mode »Adults«

If using a breathing gas humidifier, do not use an additional heat and moisture exchanger (HME)! Risk of increased breathing resistance due to condensation.

Connecting Aquapor humidifier

Prepare Aquapor according to its separate Instructions for Use.

- 1 Hang Aquapor from rail by bracket and tighten screws.
- 2 Insert elbow connector into Aquapor.
- 3 Insert the double connector into the elbow connector.
- Fill Aquapor bowl to the upper mark with distilled water.



Connection of ventilation hoses

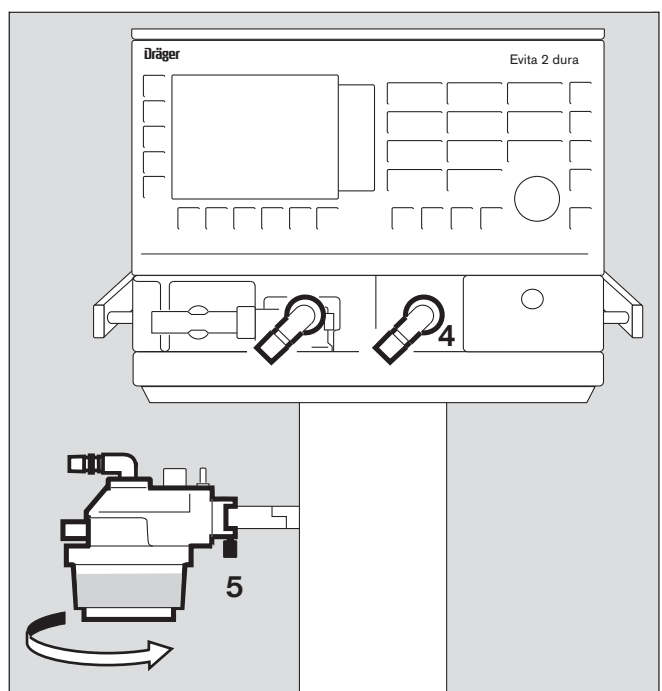
Do not use antistatic or conductive hoses*.

Depending on the desired position of the ventilator in relation to the bed, the hinged arm can be fitted to either side of the machine.

Attachment on left-hand side:

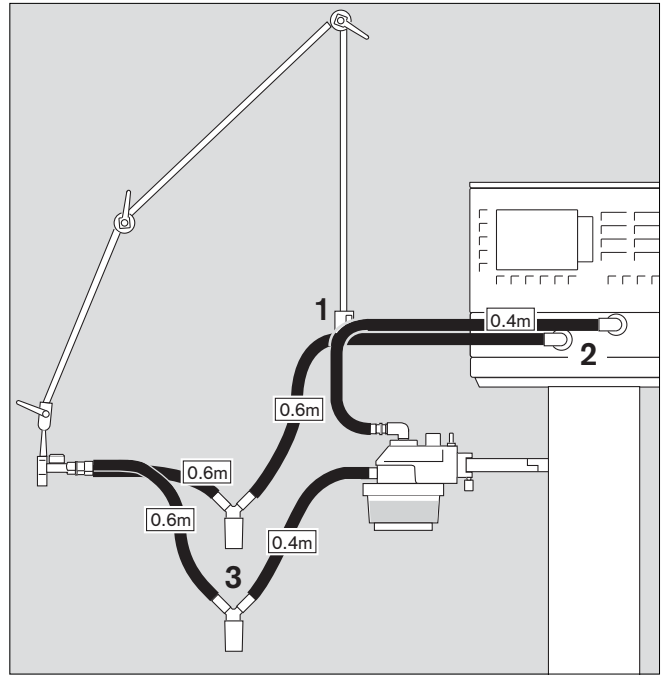
- 4 Turn both ports to the left.
- 5 Turn Aquapor to the left.

The following description applies when the ventilation hoses have been attached on the **left-hand** side.



* DIN VDE 0750 Part 215:
The use of anti-static or electrically conductive material in the breathing system of the lung ventilator is not considered to contribute any improvement in safety. On the contrary, the use of these materials increases the danger of electric shock to the patient and of fire due to the presence of oxygen.

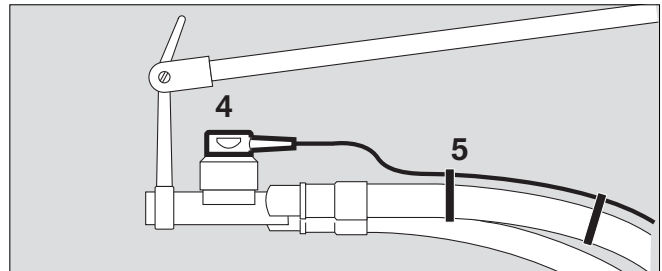
- 1 Hang the hinged arm from the rail on the left-hand side and tighten screws.
- Connect ventilation hoses, and note length of hose (metres).
- 2 Turn ports in direction of hoses.
- 3 Install water traps in vertical position at the lowest point of their hose lines.
- Connect the Y-piece, with the rubber sleeve of the Y-piece on the inspiratory side.




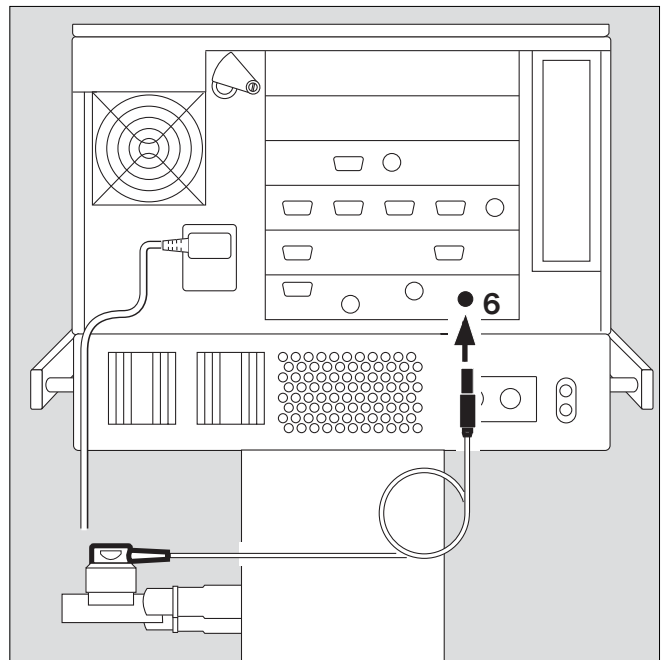
Do not place any containers with liquid on or above the ventilator! Any leaking or spilled liquid can cause malfunctions!

Fitting the temperature sensor (option)

- 4 Push the sensor as far as it will go into the rubber sleeve on the inspiratory side of the Y-piece. Align the Y-piece so that the sensor is on top.
- 5 Attach the sensor cable with hose clips.



- 6 Insert the probe of the temperature sensor into the socket »Temp.  « at the rear of the unit.



For ventilating infants

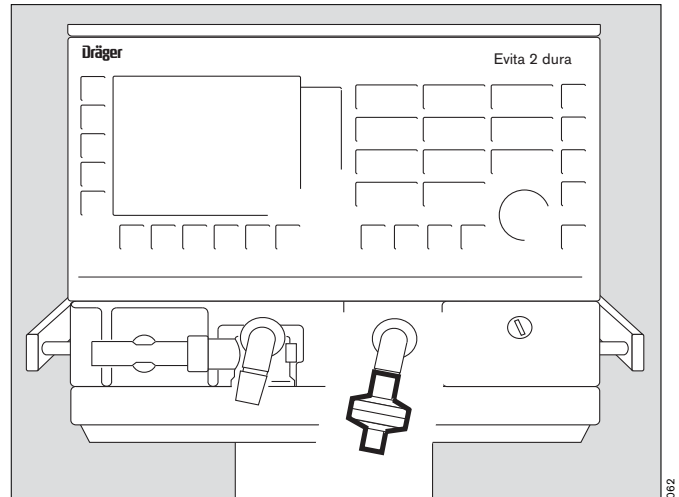
Up to 300 mL tidal volume VT and below:

Patient mode »Paediatric«

Do not use a heat and moisture exchanger (HME) at the same time as a humidifier! Risk of increased breathing resistance because of condensation.

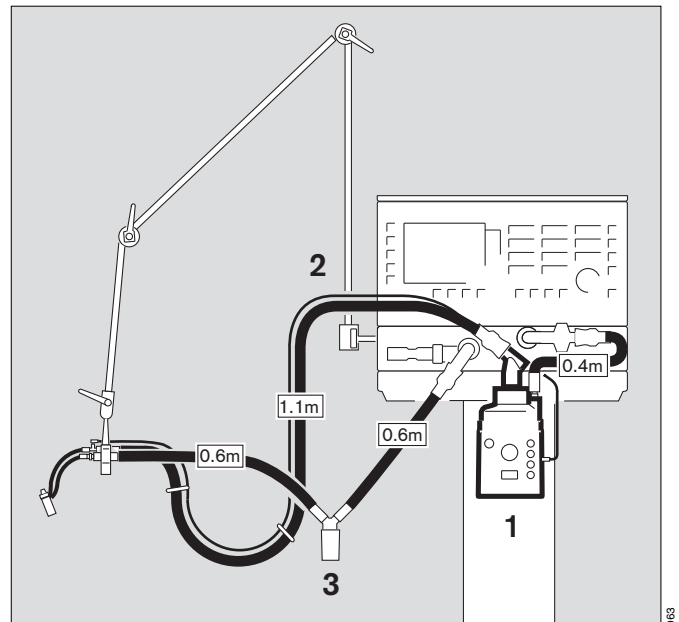
Fitting bacterial filter

- Fit the bacterial filter to the inspiratory port.



Fitting humidifier and ventilation hoses

- Prepare the "Fisher & Paykel MR 730" humidifier in accordance with the separate Instructions for Use, using hose set K (paediatric), as supplied.
- 1 Hang the humidifier with a bracket to the mounting below the machine and tighten the screws.
 - 2 Hang the hinged arm with a bracket to the rail on the left-hand side, and tighten the screws.
- Connect the ventilation hoses, and note their length (in metres).
- 3 Place the water trap in the vertical position.



Do not place any containers with liquid on or above the ventilator! Any leaking or spilled liquid can cause malfunctions!

Preparing for use

In-house/intra hospital transport

If no instrument tray (option) is fitted to the device

In-house/intra hospital transport

Ensure that the equipment does not topple over by moving the accessories as close to the ventilator and trolley as possible:

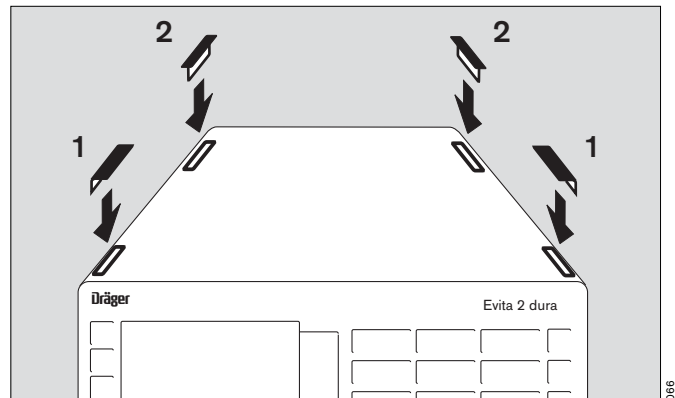
- Hinged arm set to minimum deflection.
- Drawers pushed in fully.
- Hoses and cables hooked as close as possible to the trolley.
- Humidifier secured to the trolley, not to the ceiling pendant/bed/wall.

The unit must not be tilted more than 5° when mounted on a trolley.
It may tip over at steeper angles.

If no instrument tray (option) is fitted to the device

Seal off the slits in the top panel with the rubber plugs:

- 1 Press the round plugs in the front slits – rounded part facing outwards.
- 2 Press the flat plugs into the rear slits.



Do not place any liquid container (e.g. infusion container) above or on top of the Evita 2 dura!

Any leak, spill or seepage could prevent it working properly.

Supply and Connections

Electrical power supply

The ventilator is designed for a mains voltage of

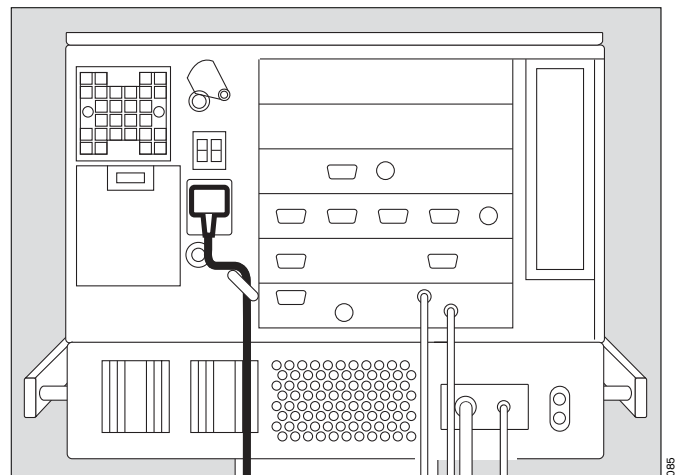
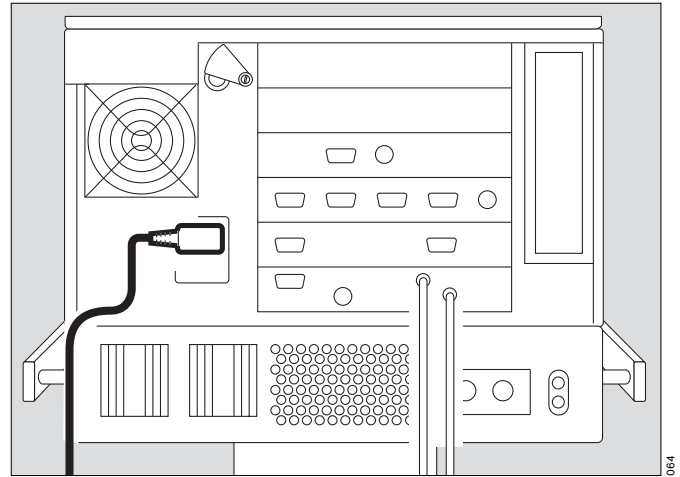
either: 220 V to 240 V

or: 100 V to 127 V

- Insert the plug in the mains socket.

Operation with Evita DC power supply and external battery (optional):

- Connect external battery via battery lead. Note Instructions for Use of the Evita DC power supply.



Note on the use of a socket strip

Connecting other devices to the same extension socket strip may, in the event of earth failure, cause the current leakage to the patient to increase beyond the permissible values. In this case, the risk of electric shock cannot be eliminated.

Temporary interruption of power supply

e.g. when switching on the reserve power supply.

Without optional Evita DC power supply:

During the power failure, Evita 2 dura will output a continuous tone for max. 2 minutes.

This continuous tone may be output for a shorter period if Evita 2 dura was switched on for less than 15 minutes.

Evita 2 dura tolerates power interruptions shorter than 10 milliseconds – without any effect on ventilation.

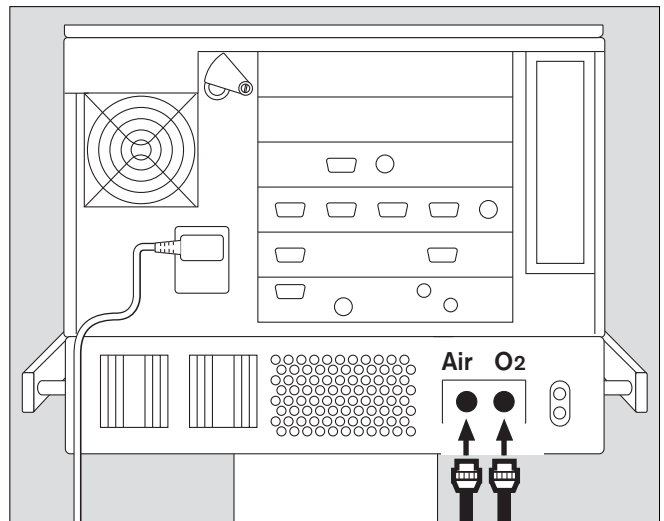
In the case of a power interruption lasting longer than 10 milliseconds, the machine restarts with a short self-test lasting about 4 seconds – ventilation is continued with the same values that were set before the power interruption. If a lower alarm limit has been set for the minute volume, the »!!! MV low« alarm is activated until the measured value has risen above the lower alarm limit.

With optional Evita DC power supply:
Follow Instructions for Use of the Evita DC power supply.

Other equipment, e.g. printers, may only be connected to the COM ports if Evita 2 dura is connected to the mains power supply via a mains power cable or if it has been earthed via the earth connection on the back of the unit. Electric power may pose a hazard in all other cases.

Gas supply

- Screw the connecting hoses for medical air and oxygen to the back panel of Evita 2 dura and insert their probes into the terminal units.
The compressed gases must be dry and free from dust and oil. Gas pressure must be 3 to 6 bar.





Evita Remote

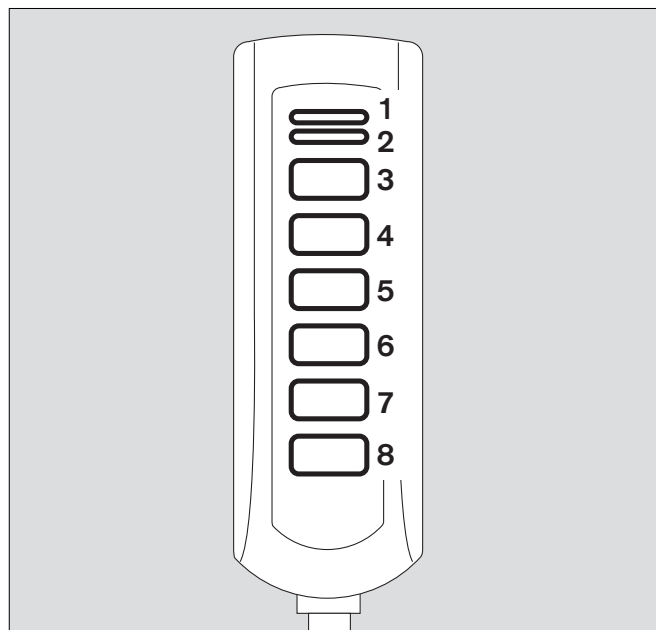
Optional remote control unit (Remote Pad)

The kit may only be installed and programmed by specialists.


For parallel, remote operation of the following LED and key functions:

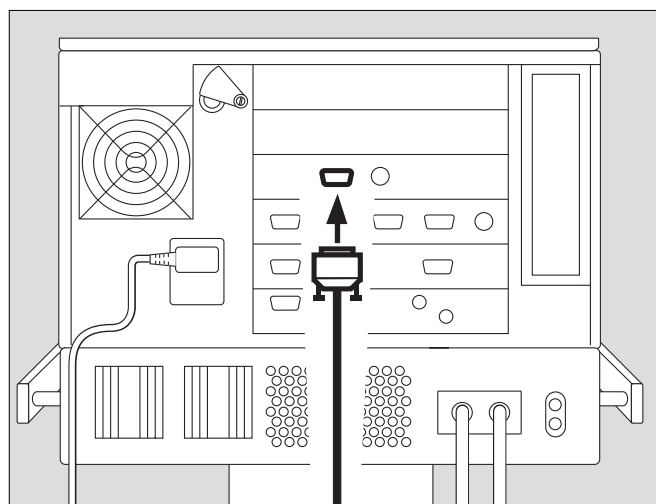
- 1 Red LED – to indicate warning messages
- 2 Yellow LED – to indicate caution and advisory messages
- 3 »« key – to suppress the alarm tone for approx. 2 minutes
- 4 »**Alarm Reset**« key – to acknowledge alarm messages
- 5 » **Neb.**« key – to start and end medicament nebulisation
- 6 »**O₂ ↑ Suction**« key – for bronchial suctioning
- 7 »**Insp. hold**« key – for sustained, manually induced inspiration
- 8 »**Exp. hold**« key – for extended and sustained expiration

The function of the respective LEDs and keys is the same as that of the corresponding elements on the front panel of Evita 2 dura and is described in the application chapters of the Instructions for Use.

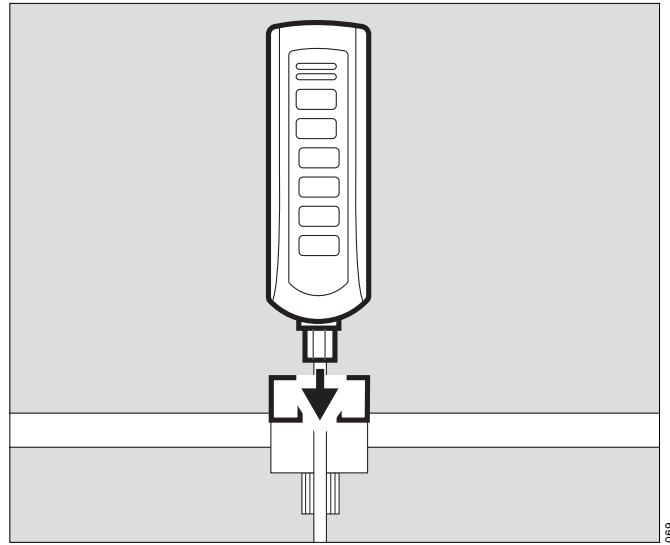


Connection

- Plug the lead of the Remote Pad into the socket »« on the rear of Evita 2 dura. The plug can be connected or disconnected at any time without impairing operation of Evita 2 dura.



- Hook holder onto a standard rail and clamp into place.
- Hang Remote Pad into holder from above.



Note automatic self-test

- when connecting the Remote Pad to Evita 2 dura while the latter is switched on
- or
- when switching on Evita 2 dura after connecting the Remote Pad.
- Do not press any keys on the Remote Pad.
- All LEDs on the Remote Pad light up for 5 seconds:
 - Red LED
 - Yellow LED
 - Yellow LEDs in the keys
- The Remote Pad is tested by Evita 2 dura. An advisory message is output if a fault is detected, see page 84 "Fault – Cause – Remedy".


Nurse call (optional)

Connection on the rear panel of Evita 2 dura for transmitting top-priority alarm signals to a central hospital alarm system.

- The kit may only be installed by specialists.
- The 6-pin round DIN plug (female connector) must be connected to the lead for the central alarm station in the hospital by a specialist.

Connection 3-5 makes and the nurse call is activated as soon as Evita 2 dura signals an alarm.

The central hospital alarm system may only be connected to the nurse call if Evita 2 dura is connected to the mains power supply via a mains power cable or if it has been earthed via the earth connection on the back of the unit. Electric power may pose a hazard in all other cases.

- Plug the connector into the »« socket on the rear and screw into place.
- Check correct operation of connected nurse call system.

Only alarm messages of the highest priority (see page 54) are transmitted via nurse call.

Alarm messages are indicated in red with three exclamation marks in the top field of the screen, see page 54. Caution and Advisory level messages are not transmitted. The nurse call is activated also when the internal loudspeaker in the ventilator is faulty.

Connection of a nurse call does not relieve staff of their duty to check the monitoring on the Evita 2 dura screen at regular intervals.

- **Screen displays must be checked regularly.**

A fault in any of the components in the link between nurse call and central hospital alarm system (e.g. in the electronics for nurse call in Evita 2 dura, in the Evita 2 dura power supply or in the alarm generator of the central hospital alarm system) may result in failure of the nurse call.

Background: The hospital connections to the central alarm typically use only one channel. The electronics for nurse call consequently also uses only one channel.

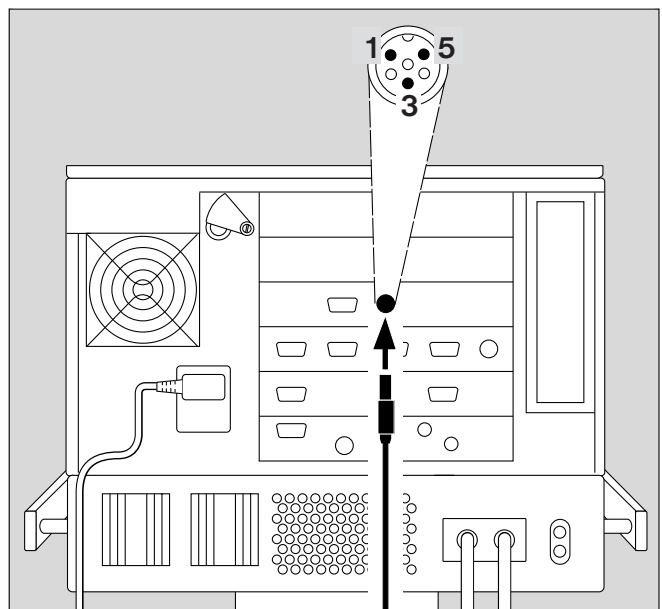
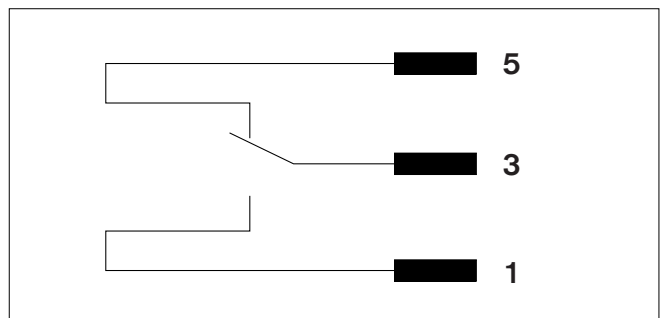
Technical Data

Floating DC contact

Input voltage Max. 40 V =

Input current Max. 500 mA

Switching capacity Max. 15 W



Device Check

Must be carried out immediately before use on the patient in order to confirm that the ventilator is operating correctly.

The device check comprises of the »Device« test and »Leakage« test.

The following functions are performed during this device check:

- Checking that the machine assembly is complete
- Testing the alarm tone
- Testing the expiratory valve
- Testing of the air-O₂ changeover valve
- Testing the safety valve
- Testing the displays
- Calibrating the flow sensor
- Calibrating the O₂ sensor
- Calibrating the CO₂ sensor
- Testing the leakproofing of the hose system
- Checking the compliance of the hose system

The test results obtained from this device check and the calibration and zero-checking values of the sensors remain stored until the next calibration – even if the device is switched off.

If the hose system, type of humidification or patient mode is changed after performing the device check, the leakproofing test must be repeated before starting operation.

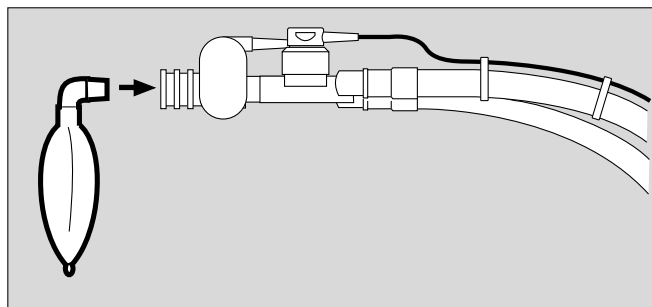
Preparing the adult test lung 84 03 201

for the adult hose system

The test lung consists of an elbow connector for connection to the Y-piece, a 7 mm diameter catheter connection for simulating the resistance of the airways and a 2 litre breathing bag to simulate compliance.

Overextended breathing bags must not be used as they may cause artefacts during the device check!

- The elbow connector must not be plugged into the patient connection of the Y-piece until directed by Evita 2 dura.

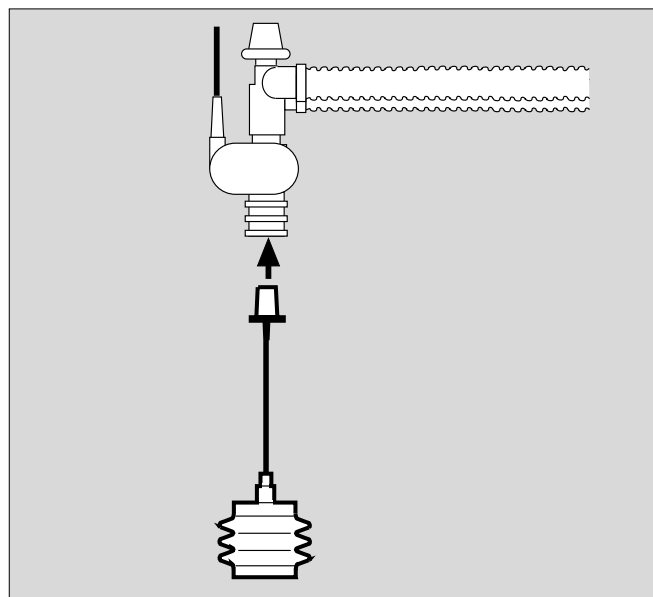


Preparing the child test lung 84 09 742

for the paediatric hose set

The test lung consists of a tracheal tube CH 12 to simulate the resistance of the airways and a small bellows to simulate compliance.

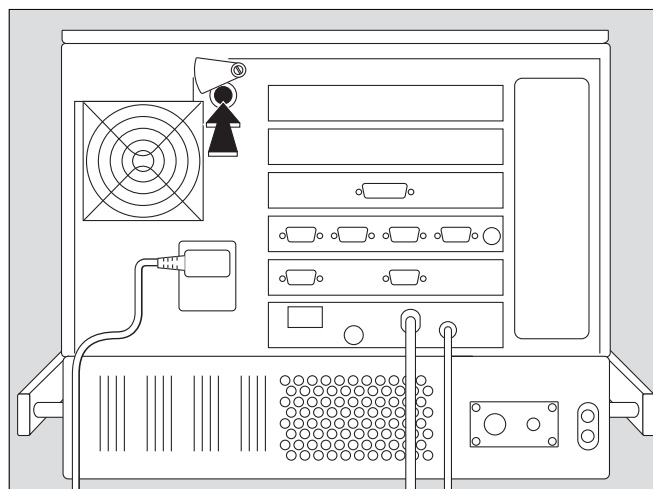
- Only insert the elbow connector into the Y-piece when Evita 2 dura advises you to do so on the screen.



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Performing the device check

- Switch on the machine = raise flap* and press power switch on the back panel until it clicks into position.
- Evita 2 dura runs through its self-test procedure.
- Wait until the 10-second test phase has been completed.

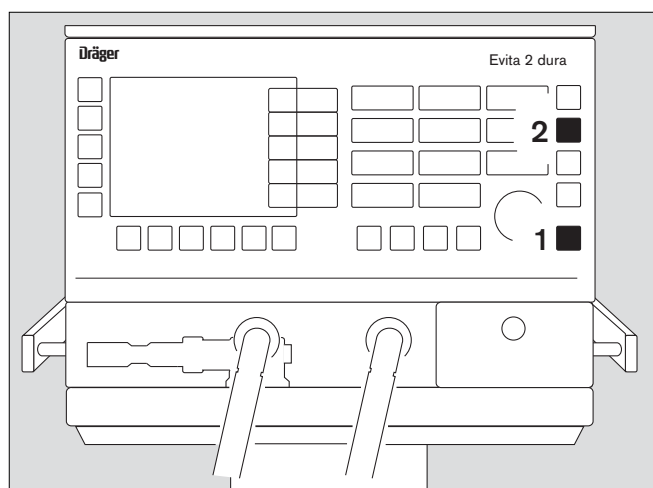


077

After the self-test:

- 1 Switch Evita 2 dura to standby = Hold down key »⏻« for about 3 seconds.
 - 2 Switch off the standby alarm tone with the »Alarm Reset« key.
- Touch the »Device check« menu key.

The standby alarm tone cannot be switched off with the »🔊« key.



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* Mains power switch flaps may differ, depending on the power supply used, see "Switching on", page 38.

Select type of humidifier

The selected type of humidifier (either active or HME) must be entered before starting the device check and leak test:

- Active humidifier, e.g. Dräger Aquapor
or
- HME/Filter (artificial nose)

If the type of humidifier is known, Evita 2 dura can take the temperature and moisture situation into account when measuring the volume parameters.

- Touch the »Humidification« screen key.
- Touch the »Active Humid.« screen key
or
- Touch the »HME/Filter« screen key = turn rotary knob.
- Confirm selection = press rotary knob.

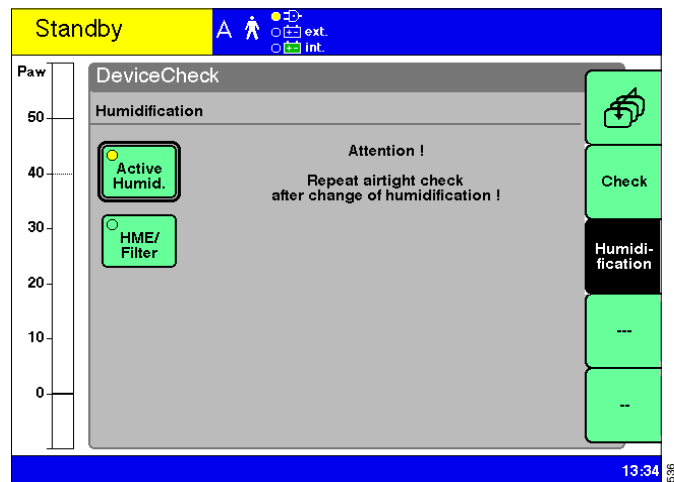
The selected humidifier is indicated by a yellow LED.

The selected type of humidifier is indicated by a black dot in the corresponding screen key.

If the type of humidifier is changed and has to be reselected on the screen, the following test steps are shown to be invalid (– – –) after the device check:

- Humidification
- Air tight check

The operator is prompted to repeat the device check for these two steps.



Start »Device« check

- Press the »Check ▶▶« menu key and select the menu »Device«.
- Activate the »Start« screen key = press rotary knob.

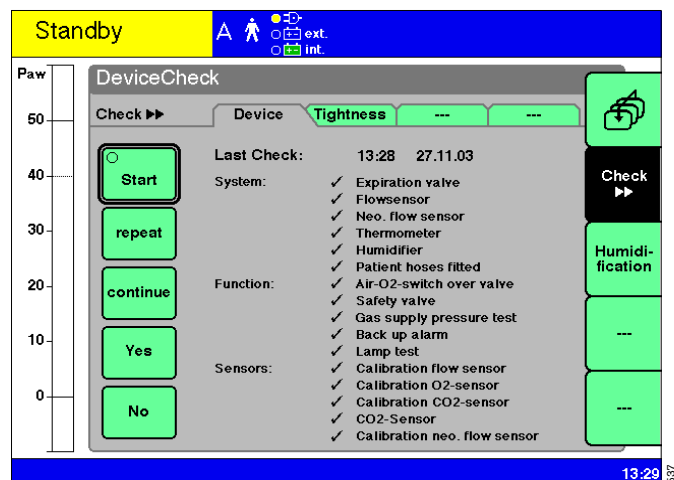
Evita 2 dura starts with the dialogue-oriented check.

The check procedure is semi automatic.

The Evita 2 dura user is instructed to carry out specific actions on the device.

A device check cannot be started during automatic calibration of the flow sensor or O₂ sensor:

- Wait until calibration is complete and then start the device check again.



The following tests are performed during the device check:

System

- Secure positioning and clear passage of the expiratory valve
- Secure positioning of the flow sensor
- Secure positioning of the neonatal flow sensor (if "NeoFlow" option is installed)
- Secure positioning of the temperature sensor
- Humidifier
- Hose system

Function

- Function of the air-O₂ changeover valve
- Function of the safety valve
- Gas supply
- Displays

Sensors

- Calibration of the flow sensor
- Calibration of the O₂ sensor
- Zero calibration of the CO₂ sensor (if "CapnoPlus" option is installed)
- CO₂ sensor (if "CapnoPlus" option is installed)
- Calibration of the neonate flow sensor (if "NeoFlow" option is installed)

The user is guided through each check in a dialogue with the device. Questions are displayed in the information line and must be answered by entering »Yes« or »No«. Instructions for carrying out the check may also be displayed instead.

On completion of the »Device« check, a checklist is displayed on the screen to show the results of the check.

Correct result: ✓
 Incorrect result: F
 Check not performed: - - -

In the event of incorrect results:

- Eliminate the cause of the fault.
- Select the »repeat« screen key = turn the rotary knob.
 Activate = press the rotary knob.

Only the tests with incorrect results are repeated.

After the »Device« check:

- Perform the leak test, page 34.

Perform »Tightness« test

- after the »Device« check
- after changing the hose system or humidifier
- after changing the patient mode
- after changing the type of humidifier
- after cancelling the leak test

The actual leakage flow is displayed continuously throughout the test. A leakage flow of 300 mL/min at a pressure of 60 mbar is permissible.

Evita 2 dura determines the compliance and resistance of the patient circuit during the »Tightness« test.

The established compliance of the hose system is used by the ventilator for automatic correction of the volume-controlled ventilation strokes, as well as of the measured values for flow monitoring, see page 123.

The established resistance of the patient circuit is used by the ventilator for correction of the pressure measured in the presence of a basic flow (NeoFlow option).

When changing the patient mode or type of humidifier, and when cancelling the leak test, the ventilator automatically resets the values for patient circuit compliance and patient circuit resistance to the default values. The »Tightness« test is shown to be invalid (– – –) and the operator is prompted to repeat the test.

- Select the menu »Tightness« via the menu button »Check ▶▶«.
- Activate screen key »Start« = press rotary knob.

The »Tightness« test can also be started separately.
In standby mode:

- Press menu key »DeviceCheck«
- Use menu key »Check ▶▶« to select the menu »Tightness«.

Leakage:

The ventilator establishes the leakage in L/min referred to a pressure of 60 mbar. Corrective measures can be taken with the aid of the continuous leakage display.

Compliance:

The ventilator establishes the system compliance in mL/mbar. The established system compliance is used for automatic correction of the volume-controlled ventilation strokes, as well as of the measured values for flow monitoring.

Resistance:

The ventilator establishes the sum of the inspiratory and expiratory resistance in mbar/L/s.



For immediate operation (e.g. in an emergency) the device check can be interrupted:

- Press the »⏻« key. The device immediately starts ventilation.

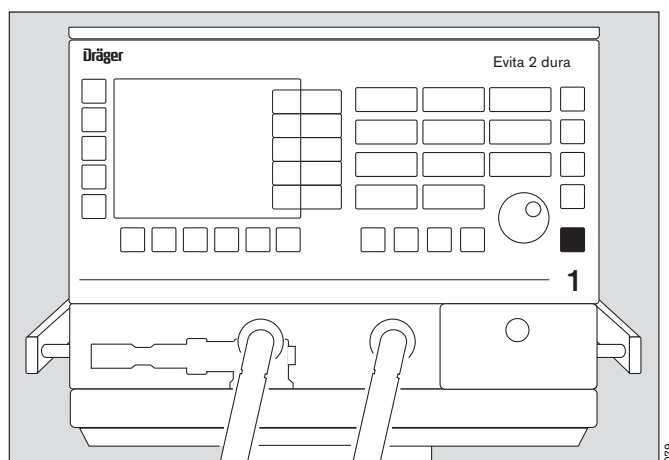
Evita 2 dura is ready for use when the device check and leak test have been successfully completed.

Either:

- Leave Evita in Standby mode and if necessary preselect the ventilation mode and ventilation parameters,

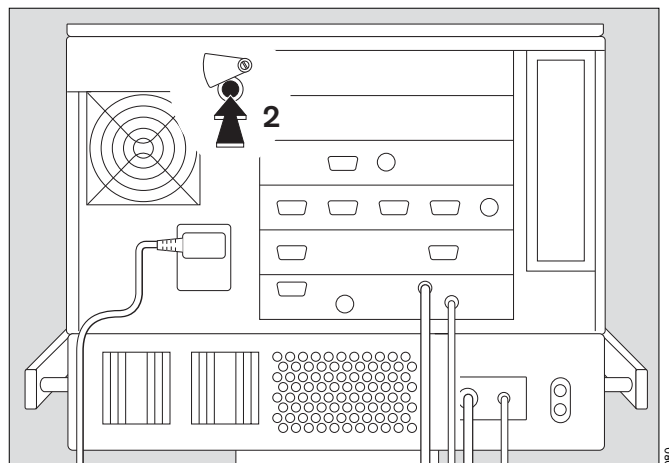
or:

- 1 Immediately start up Evita 2 dura, press »⏻«,



or:

- 2 Switch off Evita 2 dura for later use. Switch on back panel = pivot the cover-flap to the side, press button in as far as it will go and release.



Operation

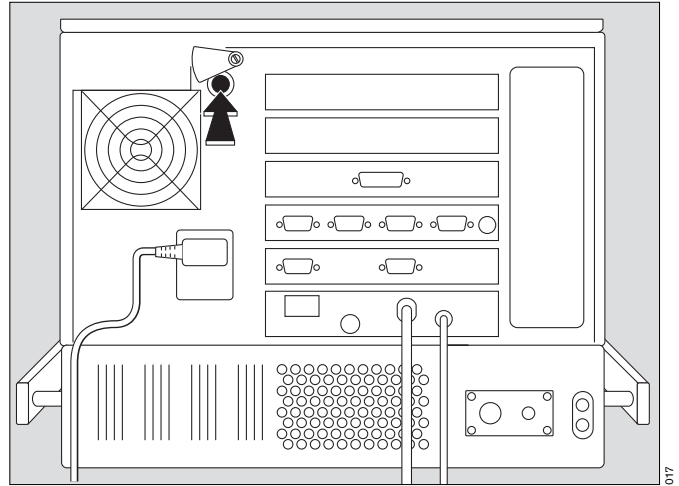
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Operation

Starting up

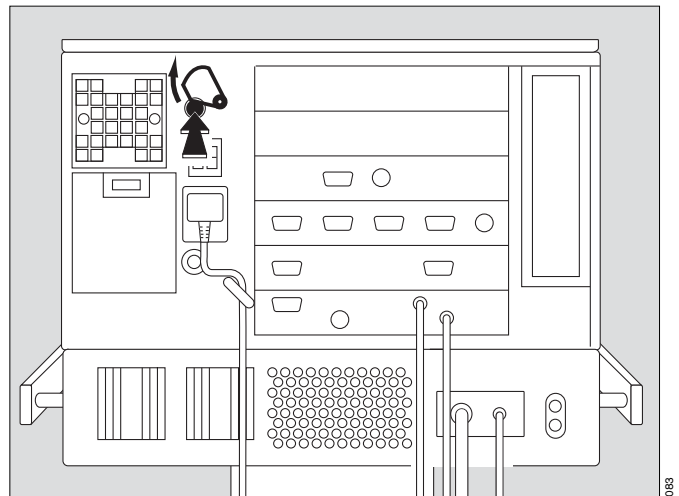
Switching on

- Push in power switch on back panel until it clicks into place = ON.
The flap comes down over the switch to prevent it being inadvertently switched off.



Units with DC power supply MB:

- Switch on unit = pivot flap upwards and press power switch on rear of unit until it clicks into position. The flap falls over the button to protect against inadvertent switching off. To switch the ventilator off, first pivot the mains power switch flap upwards and then depress the button fully, then release the button and allow the flap to settle back into position.



Evita 2 dura runs a self-test.

- Wait until the 10-second test phase is complete.
- Evita 2 dura always begins ventilation with the start-up values marked by an arrow on the on-screen knobs.
- To select these start-up values, please refer to pages 77 onwards.
- After power cuts and after standby mode, the settings valid immediately before the interruption of operation remain in use.

Patient mode

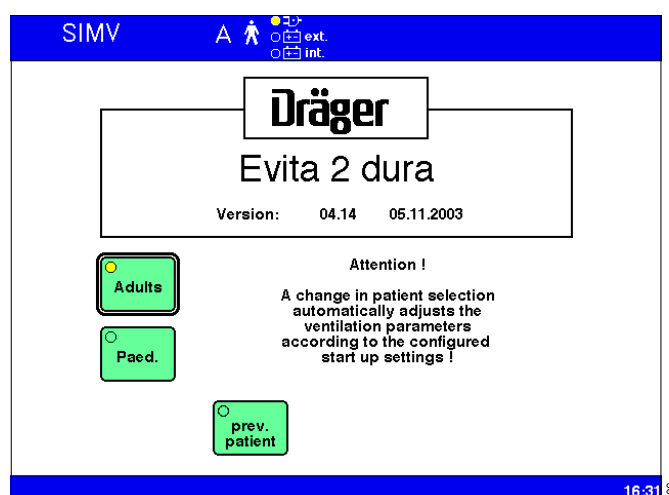
The patient mode can be selected after switching on Evita 2 dura:

- »Adults« = adult patients
- »Paed.« = children
- »Neo.« = neonates (when using the "NeoFlow" option)
- »prev. patient« = previous patient

Example:

Adult ventilation

With this information, Evita 2 dura defines the adjustment ranges and the start-up values of the ventilation parameters.

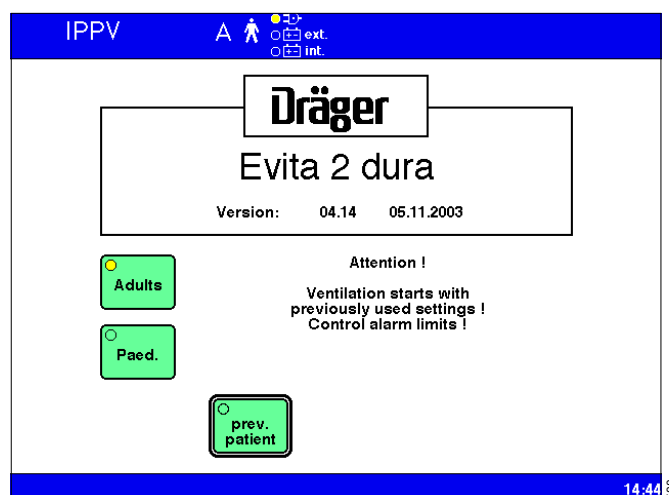


The screen key »prev. patient« can be used to restore the specific patient settings, including alarm limits and monitoring status, effective before switching off the device.

The previous modes are displayed in the status line:

- Previous ventilation mode (example: IPPV)
- Previous patient mode (A = Adult)
- Previous application mode – tube or mask for optional NIV (example: NIV)

The key »prev. patient« is not displayed by Evita 2 dura following a loss of data or removal of a previously used option (e.g. NeoFlow), thus preventing restoration of the previous setting.



Selecting the patient mode

Either:

- Select the »Adults« key or the »Paed.« key or the »Neo.« key (NeoFlow option) = turn rotary knob.
- Confirm = press rotary knob.

or:

- Select the key »prev. patient« = turn rotary knob.
- Confirm = press rotary knob.

Starting ventilation

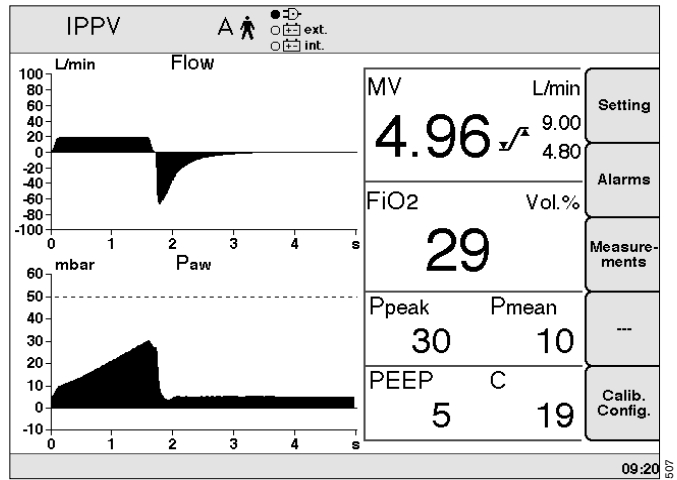
Evita 2 dura starts ventilation with the ventilation mode configured by the user and with the specific start settings for the patient or with the settings valid before the machine was last switched off.

To select other start-up settings, see page 77.

If no selection is made or if the rotary knob is not pressed to confirm the new settings, the apparatus automatically starts ventilation after 30 seconds with the last selected patient mode and ventilation mode and the associated ventilation parameters.

The main page is displayed on the Evita 2 dura screen.

The user can check and correct the settings in the display fields next to the parameter keys.

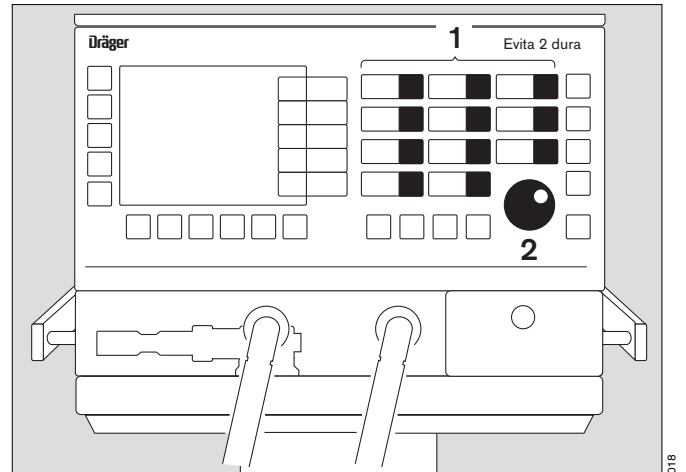


Setting Ventilation Modes

To set the ventilation parameters:

- 1 Press the appropriate ventilation mode key. The yellow LED in the key will light up.
- 2 Set the desired value = turn the rotary knob.
Confirm value = press the rotary knob. The yellow LED will go out.
If the setting is at the upper or lower limit of the adjustment range for a parameter, the LED in the relevant key will start flashing.
- 2 Acknowledge = press the rotary knob.

If you fail to confirm/acknowledge the new settings within 30 seconds, the previous settings will remain operative.



Setting parameters for another ventilation mode

- 1 Press the relevant ventilation mode key briefly.
Its LED will flash.
The yellow indicators for the relevant parameters for the new ventilation mode flash in the parameter keys.

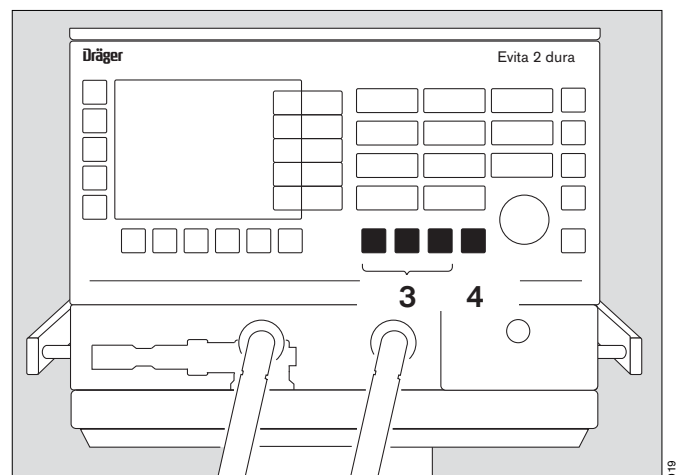
Set the new ventilation parameters:

- 1 Press the relevant key: its LED will stop flashing and remain constantly lit.
- 2 Set the desired value = turn the rotary knob.
Confirm value = press the rotary knob. The yellow LED will go out.


To activate the ventilation mode:

- 3 Ventilation mode keys:
 - IPPV
 - SIMV
 - BIPAP
 or
- 4 »Other modes« key for other ventilation modes that are set on the screen.
Factory-set default: CPAP/ASB.
- Hold down the relevant key for 3 seconds,
or
- briefly press and release the relevant key and press the rotary knob.


The selected ventilation mode will now be active.

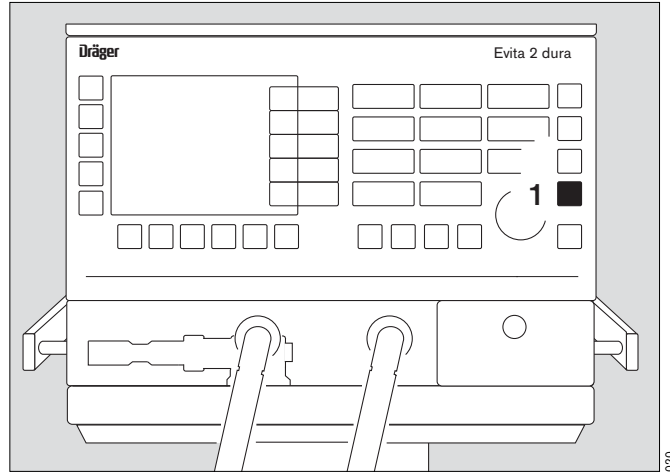


To prevent settings being changed inadvertently:

- 1 Press the »**1**« key. Its yellow LED will light up. The parameter keys and ventilation mode keys will be protected against inadvertent setting.

Before setting a new value:

- 1 Press the »**1**« key. The yellow LED will go out.



IPPV

Intermittent Positive Pressure Ventilation

Volume-controlled ventilation with a fixed mandatory minute volume MV, set with the tidal volume V_T and frequency f .

For patients unable to breathe spontaneously.

Set ventilation pattern for IPPV via the keys for the ventilation parameters:

Tidal volume »**VT**«

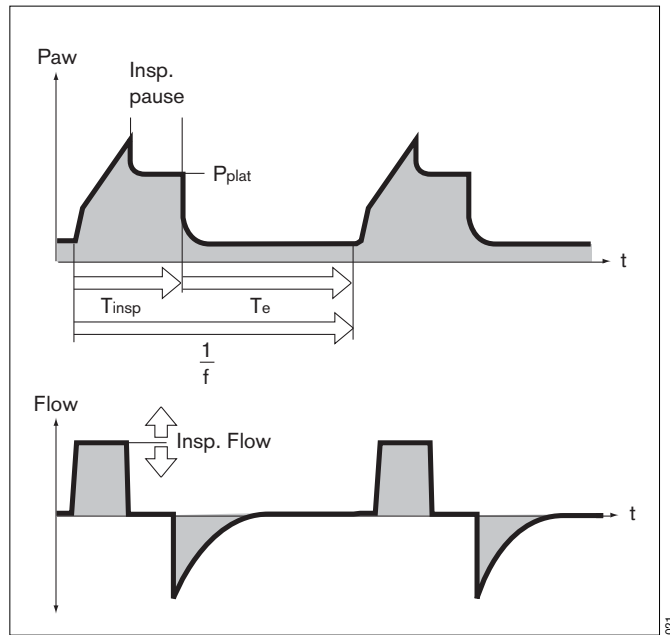
Insp. Flow »**Flow**«

Frequency »**f**«

Inspiration time »**T_{insp}**«

O₂ concentration »**O₂**«

Positive end-expiratory pressure »**PEEP**«



IPPV can be supplemented by the following ventilation parameters:

Trigger (IPPV_{Assist}) – for synchronising mandatory ventilation with attempted spontaneous breathing by the patient. By activating the trigger and setting the trigger sensitivity, the mandatory ventilator strokes are synchronised with the patient's spontaneous breathing attempts.

The trigger can be switched off if synchronisation with the patient's spontaneous breathing attempts is not required.

To activate/set:

- Press ventilation parameter key »**Trigger**«.
- Set value = turn the rotary knob,
Confirm value = press the rotary knob.

To deactivate:

- Set a value less than 0.3 or above 15 L/min.
The display will show: – – –

Sigh – to prevent atelectasis.

Atelectasis can be prevented by activating the Sigh function and setting the sigh in the form of an intermittent PEEP. When the Sigh function is activated, the end-expiratory pressure is increased by the set intermittent PEEP for 2 ventilation strokes every 3 minutes.

Pmax

IPPV can be supplemented by the ventilation parameter Pmax.

- Activate »Pressure limit Pmax«, see page 79.
- Set value Pmax via the key for the ventilation parameter »Pinsp«.

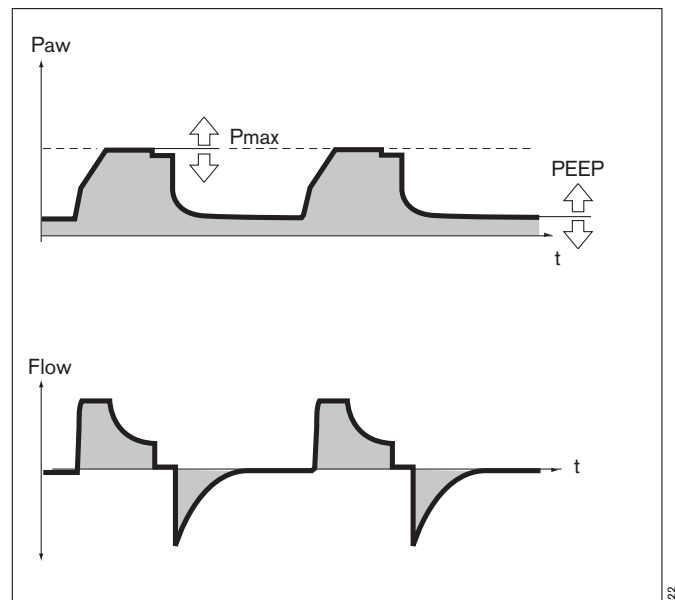
Pressure limited ventilation PLV* – for manually limiting pressure peaks to the pressure limit Pmax.

The tidal volume remains constant as long as the pressure curve continues to show a short pressure plateau and the flow curve shows a brief pause in the flow between inspiration and expiration.

Volume monitoring is constantly active. If the set tidal volume V_T can no longer be applied, the alarm »!! Volume not constant« is automatically generated.

When changing from IPPV to a pressure-controlled ventilation mode, the value of Pmax is adopted but is limited to 50 mbar (the display for the »Pinsp« ventilation parameter flashes).

- Confirm value = press the rotary knob
or
set a higher value.



* For a detailed description of PLV, see page 116.

SIMV, SIMV/ASB

**Synchronized Intermittent Mandatory Ventilation*
Assisted Spontaneous Breathing****

Fixed mandatory minute volume MV, set with the tidal volume VT and frequency f. The patient can breathe spontaneously between the mandatory ventilation strokes, thereby contributing to the overall minute volume. Spontaneous breathing can be assisted by ASB.

For patients with insufficient spontaneous breathing or patients being weaned from artificial ventilation by progressive reduction of the mandatory proportion of the total minute volume.

The frequency can be reduced to 0 during the weaning process. The machine automatically changes to ventilation mode CPAP or CPAP/ASB. This ventilation mode is also displayed.

Set the ventilation pattern for SIMV via the keys for the ventilation parameters:

- Tidal volume »VT«
- Insp. Flow »Flow«
- Frequency »f«
- At f = 0/min, the ventilator switches to CPAP mode.
- Inspiration time »T_{insp}«
- Sensitivity »Trigger«
- O₂ concentration »O₂«
- Positive end-expiratory pressure »PEEP«

Additionally for SIMV/ASB:

- Pressure support »PASB«
- Pressure rise time »Ramp«

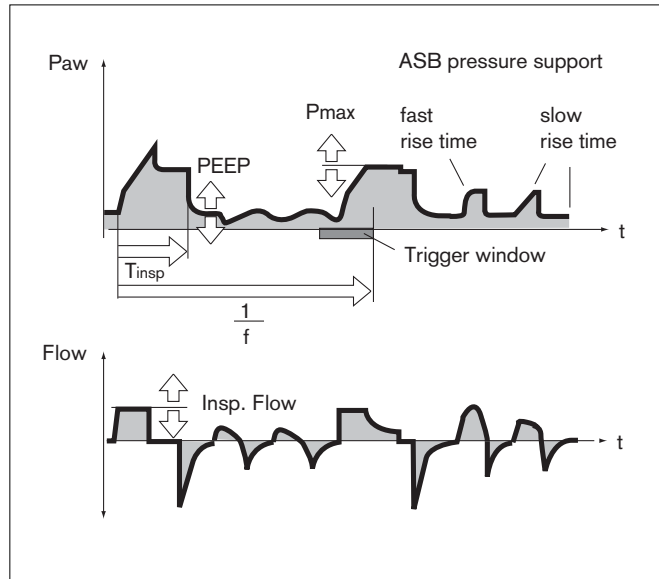
SIMV, SIMV/ASB can be supplemented with the following ventilation parameters:

Apnoea ventilation – for automatic switchover to volume-controlled mandatory ventilation if the patient stops breathing. If breathing stops, Evita 2 dura activates an alarm after the set alarm time (T_{Apnoea} /^f) and starts volume-controlled ventilation with the set ventilation parameters:

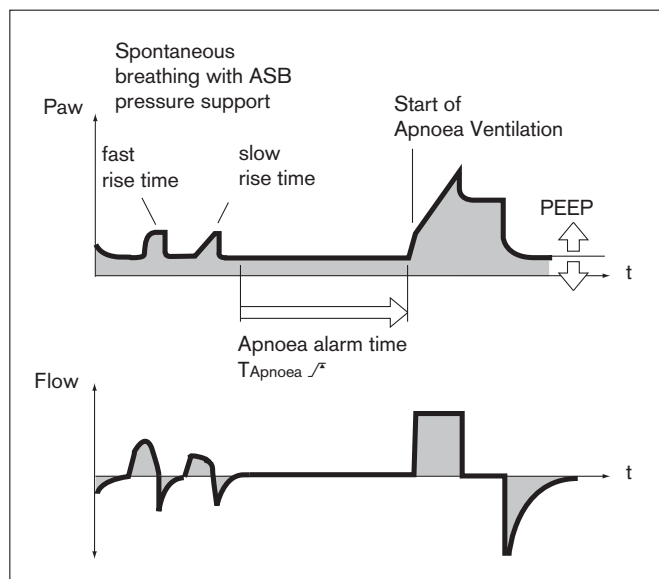
- Frequency »f_{Apnoea}«
- Tidal volume »VT_{Apnoea}«

The patient can breathe spontaneously during apnoea ventilation. The apnoea ventilation frequency remains constant.

Set apnoea ventilation, see page 52.



* For a detailed description of SIMV, see page 118.
** For a detailed description of ASB, see page 119.



Pmax

SIMV, SIMV/ASB can be supplemented with the ventilation parameter Pmax:

- Activate »Pmax pressure limit«, see page 79.
- Set the value of Pmax with the »P_{insp}«

Pressure limited ventilation PLV* – for manually limiting pressure peaks using the Pmax pressure limit.

The tidal volume remains constant as long as the pressure curve continues to show a short plateau, and the flow curve shows a brief pause in the flow between inspiration and expiration.

Volume monitoring is constantly active. If the set tidal volume VT can no longer be applied, the alarm »!! **Volume not constant**« is automatically generated.

* For a detailed description of PLV, see page 116.

BIPAP, BIPAP/ASB

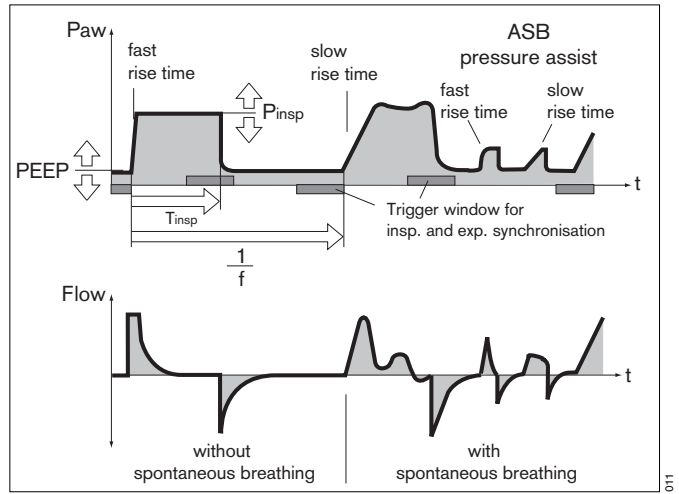
Biphasic Positive Airway Pressure Assisted Spontaneous Breathing

Pressure-controlled ventilation combined with free spontaneous breathing during the complete breathing cycle, supported by adjustable additional pressure at CPAP level.

The mandatory proportion of the total minute volume MV is set by means of the inspiration pressure P_{insp} , PEEP and frequency f .

Adaptable to a wide range of patients, from those unable to breathe spontaneously at all to those breathing spontaneously before extubation. Suitable for weaning patients from artificial ventilation by progressively reducing the mandatory fraction of the minute volume MV and by reducing the additional artificial pressure support PASB.

The frequency can be reduced to 0 during the weaning process. The machine automatically changes to ventilation mode CPAP or CPAP/ASB. This ventilation mode is also displayed.



Set the ventilation pattern for BIPAP via the keys for the ventilation parameters:

Inspiration pressure »**P_{insp}**«

If P_{insp} is set to the same value as PEEP, the machine changes over to CPAP mode.

Frequency »**f**«

If $f = 0/\text{min}$, the apparatus switches over to CPAP mode.

Inspiration time »**T_{insp}**«

Sensitivity »**Trigger**«

O₂ concentration »**O₂**«

Positive end-expiratory pressure »**PEEP**«

Additionally for BIPAP/ASB:

Pressure support »**PASB**«

Pressure rise time »**Ramp**«

BIPAP, BIPAP/ASB can be supplemented with the following ventilation parameters:

Apnoea ventilation – for automatic switchover to volume-controlled mandatory ventilation if the patient stops breathing.

If the patient stops breathing, Evita 2 dura activates an alarm after the set alarm time ($T_{Apnoea} \sqrt{\wedge}$) and starts volume-controlled ventilation with the set ventilation parameters:

Frequency »**f_{Apnoea}**«

Tidal volume »**V_{TApnoea}**«

The patient can breathe spontaneously during apnoea ventilation. The apnoea ventilation frequency remains constant.

Set apnoea ventilation, see page 52.

BIPAP_{Assist}**Biphasic Positive Airway Pressure Assisted**
Druckkontrollierte, assistierende Beatmung

The inspiratory strokes are the same as for BIPAP, but the changeover from P_{insp} to PEEP is not synchronised with expiration by the patient. The patient can breathe spontaneously at PEEP level through the entire ventilation process.

Every spontaneous breathing activity by the patient triggers a synchronised inspiratory stroke.

A non-synchronised inspiratory stroke is started by the device at the latest upon expiry of the time $\frac{1}{f}$.

For all patients, from those unable to breathe spontaneously to those breathing spontaneously before being weaned off the ventilator.

The set values for the relevant ventilation parameters are displayed alongside the keys for the ventilation parameters.

Set the ventilation pattern for BIPAP_{Assist} via the keys for the ventilation parameters:

Inspiratory pressure »**P_{insp}**«

Frequency »**f**«

Inspiration time »**T_{insp}**«

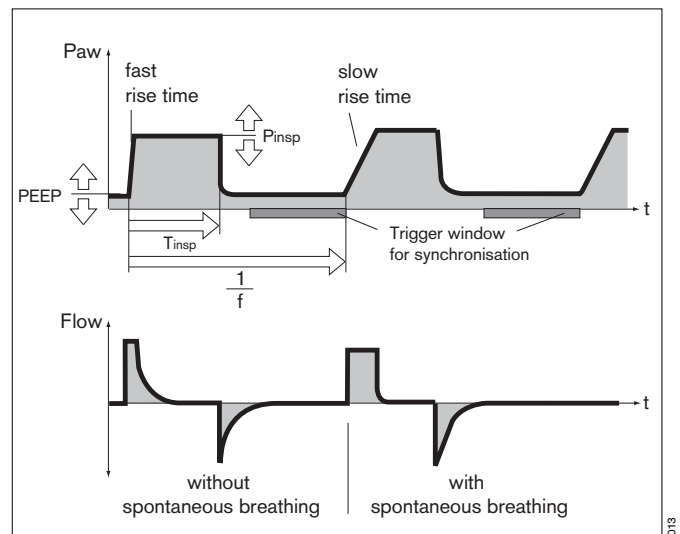
Sensitivity »**Trigger**«

O₂ concentration »**O₂**«

Positive end-expiratory pressure »**PEEP**«

Pressure rise time »**Ramp**«

Absolute inspiratory pressure »**P_{insp}**«



To activate:

- Press menu button »**other modes**«.

When BIPAP_{Assist} has been selected in the menu with the cursor:

- Hold the menu button »**other modes**« for approx. 3 seconds,

or:

- Press the rotary knob.

Otherwise:

- Select BIPAP_{Assist} = turn rotary knob, activate = press rotary knob.

The "LED" in the screen key »**BIPAP_{Assist}**« changes from green to yellow. Ventilation mode BIPAP_{Assist} is now active and displayed in the status line.

CPAP, CPAP/ASB

Continuous Positive Airway Pressure Assisted Spontaneous Breathing

Spontaneous breathing at a raised pressure level, to increase the functional residual capacity FRC. Spontaneous breathing can be assisted with additional pressure by ASB.

For patients breathing spontaneously.

Set the ventilation pattern for CPAP via the keys for the ventilation parameters:

O₂ concentration »O₂«

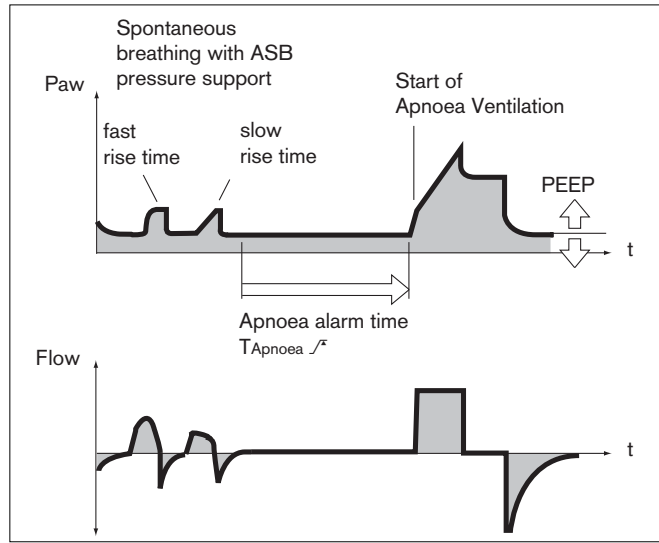
Positive end-expiratory pressure »PEEP«

Additionally, for CPAP/ASB:

Pressure support »PASB«

Pressure rise time »Ramp«

Sensitivity »Trigger«



To activate:

- Press menu button »other modes«.

When CPAP/ASB has been selected in the menu with the cursor:

- Hold the menu button »other modes« for approx. 3 seconds,

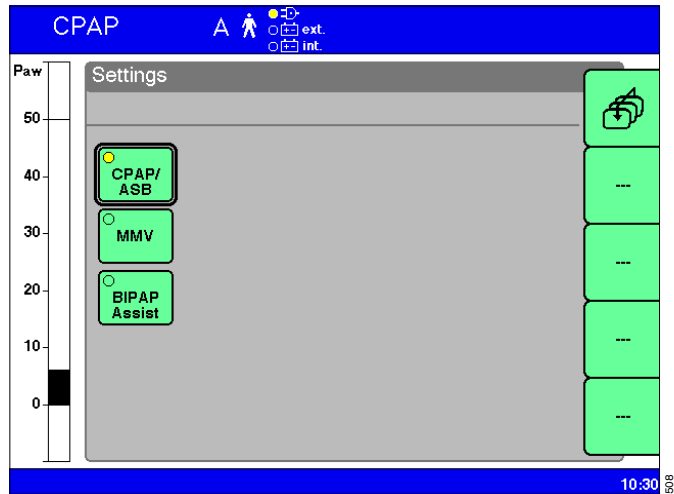
or:

- Press the rotary knob.

Otherwise:

- Select CPAP/ASB = turn rotary knob, activate = press rotary knob.

The "LED" in the screen key »CPAP/ASB« changes from green to yellow. Ventilation mode CPAP/ASB is now active and displayed in the status line.



CPAP/ASB can also be activated when frequency f = 0 in SIMV or BIPAP.

CPAP, CPAP/ASB can be expanded with the following ventilation parameters:

Trigger – for synchronising ventilation with attempted spontaneous breathing by the patient.

By activating the trigger and setting the trigger sensitivity, the assisting ventilator strokes are synchronised with the patient's own spontaneous breathing attempts.

Apnoea ventilation – for automatic switchover to volume-controlled mandatory ventilation if the patient stops breathing.

If the patient stops breathing, Evita 2 dura activates an alarm after the set alarm time (T_{Apnoea} / \bar{x}) and starts volume-controlled ventilation with the set ventilation parameters:

Frequency » f_{Apnoea} «

Tidal volume » VT_{Apnoea} «

The patient can breathe spontaneously during apnoea ventilation. The apnoea ventilation frequency remains constant.

To set apnoea ventilation, see page 52.

MMV, MMV/ASB

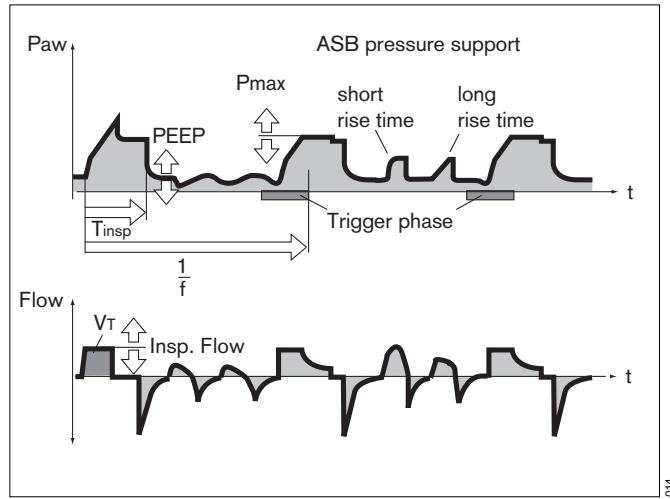
**Mandatory Minute Volume Ventilation*
Assisted Spontaneous Breathing**

The overall minute volume is preset to a mandatory level, which can be adjusted by means of the tidal volume V_T and frequency f .

The patient can breathe spontaneously, thereby contributing a proportion of the total minute volume.

The difference between the spontaneously breathed minute volume and the set minute volume is covered by the mandatory ventilation strokes. Spontaneous breathing can be assisted by the ASB pressure support.

This mode is intended for patients being weaned off the ventilator by progressively reducing the mandatory proportion of the total minute volume.



Set the pattern of ventilation for MMV, MMV/ASB with the ventilation parameters:

Tidal volume » **V_T** «

Insp. Flow »**Flow**«

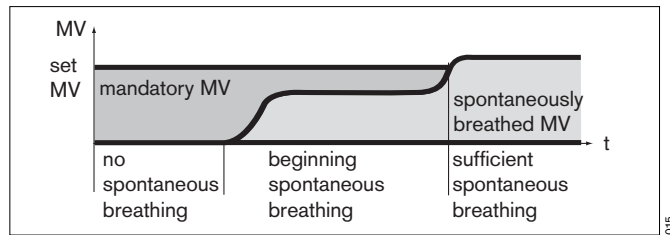
Frequency » **f** «

Inspiration time » **T_{insp}** «

Sensitivity »**Trigger**«

O₂ concentration »**O₂**«

Positive end-expiratory pressure »**PEEP**«



Additionally, for MMV/ASB:

Pressure assist »**PASB**«

Pressure rise time »**Ramp**«

To activate:

- Press menu button »**other modes**«.

When MMV has been selected in the menu with the cursor:

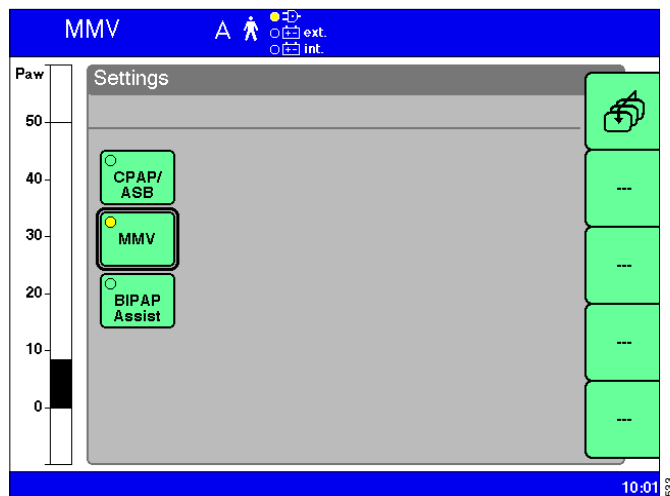
- Hold the menu button »**other modes**« for approx. 3 seconds,
- or:

- Press the rotary knob.

Otherwise:

- Select MMV = turn rotary knob, activate = press rotary knob.

The "LED" in the screen key »**MMV**« changes from green to yellow. Ventilation mode MMV is now active and displayed in the status line.



* For a detailed description of MMV, see page 122.

Pmax

MMV, MMV/ASB can be supplemented with the ventilation parameter Pmax.

- To activate »Pmax pressure limit«, see page 79.

Pressure limited ventilation PLV* – for manually limiting pressure peaks using the Pmax pressure limit.

The tidal volume remains constant as long as the pressure curve continues to show a short plateau, and the flow curve shows a brief pause in the flow between inspiration and expiration.

Volume monitoring is constantly active. If the set tidal volume VT can no longer be applied, the alarm »!! Volume not constant« is automatically generated.

* For a detailed description of PLV, see page 116.

Apnoea ventilation

For automatic switch-over to volume-controlled mandatory ventilation if the patient stops breathing.

It can be switched on in the ventilation modes SIMV, BIPAP, CPAP, APRV.

Evita 2 dura emits an apnoea alarm if during the set alarm period »T_{Apnoea}« no expiration flow is measured or insufficient inspiratory gas is delivered.

If breathing stops, Evita 2 dura emits an alarm after the set alarm time (T_{Apnoea} / √) and starts volume-controlled ventilation with the set ventilation parameters:

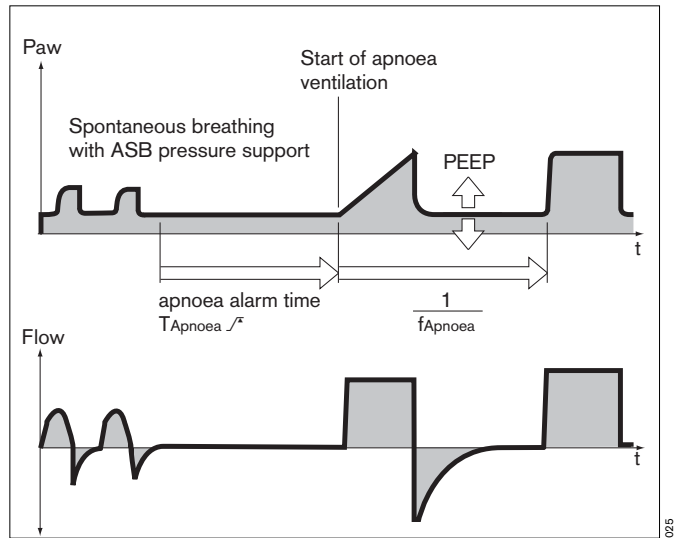
Frequency »f_{Apnoea}«

Tidal volume »V_{TApnoea}«

The ventilation parameters »O₂« and »PEEP« correspond to the settings effective at the time.

The inspiration time for apnoea ventilation is determined from the set apnoea frequency »f_{Apnoea}« and a fixed I:E ratio of 1:2.

As in SIMV, the patient can breathe spontaneously during apnoea ventilation and the mandatory ventilation strokes will be synchronised with the patient's spontaneous breathing. The apnoea ventilation frequency remains constant.

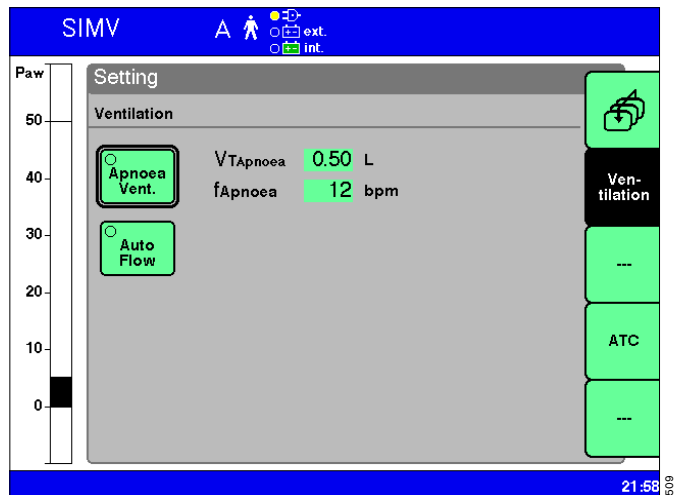


To set apnoea ventilation:

- Press menu button »Settings«.

Display:

- Select screen key »Apnoea Vent.« = turn rotary knob. Switch on apnoea ventilation = press rotary knob. The "LED" in the screen key changes from green to yellow = apnoea ventilation is on.
- Select field »V_{TApnoea}« = turn rotary knob, activate = press rotary knob.
- Set values = turn rotary knob, confirm = press rotary knob.
- Select, set and confirm »f_{Apnoea}« accordingly.



To terminate apnoea ventilation:

- Press »Alarm Reset« key. The machine will continue operating in its previous ventilation mode.
or
select another ventilation mode.

See page 80 for configuration of the apnoea ventilation status when starting the machine.

Setting Alarm Limits

- Press the »Alarms« menu key.

Example display »Limits«

This page displays all the alarm limits that can be set/adjusted.

√ = lower alarm limit

∧ = upper alarm limit

Example: Setting the upper alarm limit for fspont.

- Select the »fspont.« screen field with the cursor = turn the rotary knob.
Confirm = press the rotary knob.
- Set the desired value = turn the rotary knob.
Confirm = press the rotary knob.

The lower alarm limit does not have to be set for the airway pressure Paw, because it is automatically coupled with the PEEP setting.

The alarm limits do not have to be set for the O₂ concentration. These limits are automatically coupled to the O₂ concentration setting.

Lower alarm limit:

or settings up to 60 Vol.% O₂: setting -4 Vol.% O₂

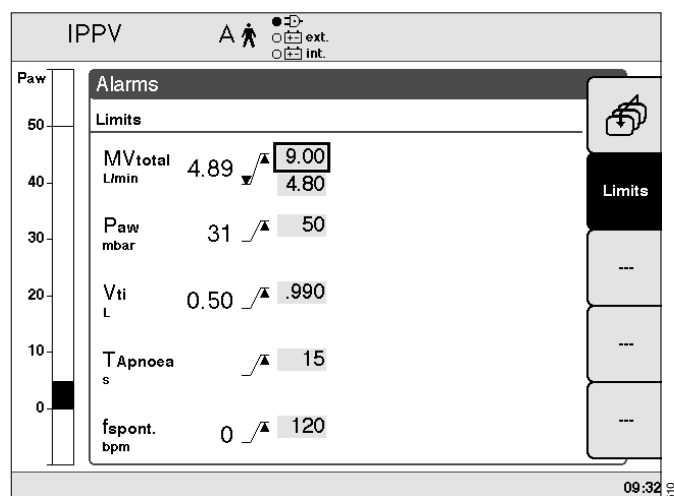
for settings from 60 to 100 Vol.% O₂: setting -6 Vol.% O₂

Upper alarm limit:

for settings up to 60 Vol.% O₂: setting +4 Vol.% O₂

for settings from 60 to 100 Vol.% O₂: setting +6 Vol.% O₂

Setting ranges for alarm limits, see "Technical Data", page 112.



In the Event of an Alarm

- 1 the red or yellow LED flashes.
- 2 The alarm message is displayed in the right-hand corner of the top line of the screen.

Evita 2 dura assesses the alarm message according to its priority, marks the text with exclamation marks and generates the various alarm tone sequences.

Warning = top priority message

- 1 The red LED flashes.

Warning messages are marked with three exclamation marks.

Example: »!!! **Apnoea**«

Evita 2 dura generates a five-tone sequence that is sounded twice and repeated every 7 seconds.

Caution = medium priority message

- 3 The yellow LED flashes.

Caution messages are marked with two exclamation marks.

Example: »!! **Check settings**«

Evita 2 dura generates a 3-tone sequence that is repeated every 20 seconds.

Advisory = low priority message

- 3 The yellow LED lights up and remains constantly lit.

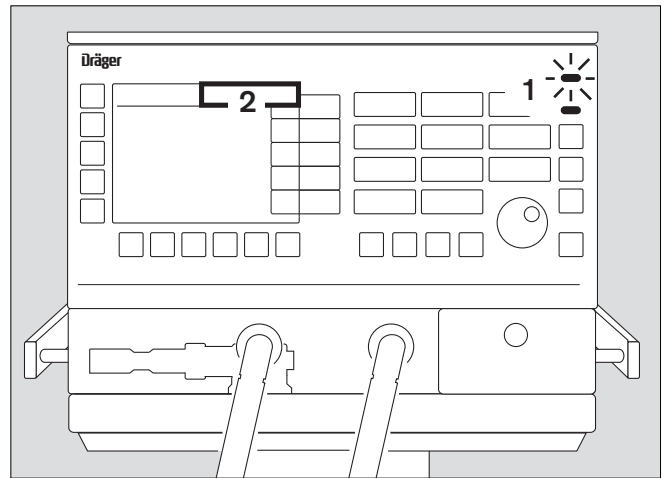
Caution messages are marked with one exclamation mark.

Example: »! **Malfunction fan**«

Evita 2 dura generates a 2-tone sequence that sounds only once.

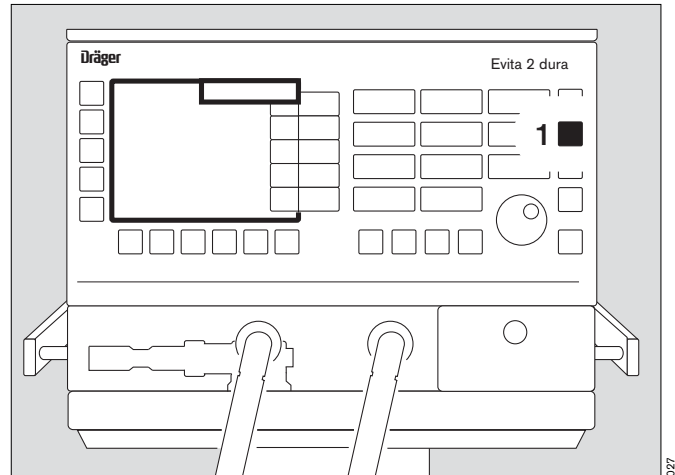
If the loudspeaker for audible alarms fails on account of a defect, an auxiliary signal will sound continuously. This continuous tone also serves as power failure alarm, see page 25, if power is interrupted while the ventilator is in use.

To remedy the faults, please refer to the "Troubleshooting" section (starting) on page 84.



Once the fault has been remedied, the alarm tone is switched off. Caution and advisory messages disappear automatically. Alarm messages (!!!) are then displayed in the colour of the status line and must be acknowledged:

- 1 Press the »Alarm Reset« key.
The message is erased from the screen.



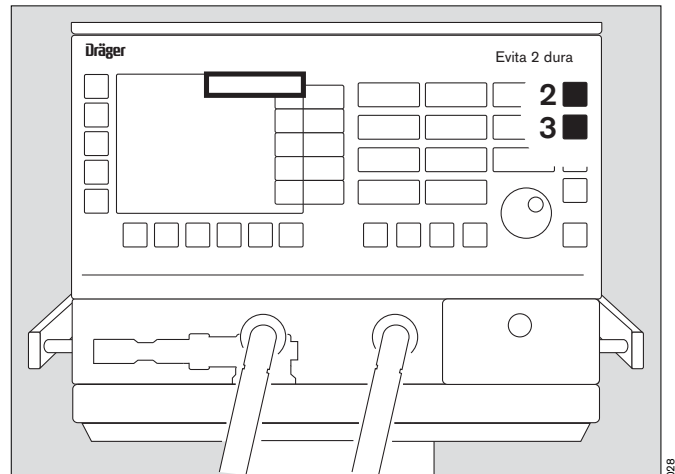
Suppressing the audible alarm

for max. 2 minutes:

- 2 Press the »Mute« key, its yellow LED lights up and the acoustic alarm is cancelled for 2 minutes.
If the fault that triggered the alarm is still not remedied, the audible alarm starts up again after this period.

If you wish to reactivate the audible alarm before the end of the 2-minute muting period:

- 2 Press the »Mute« key again, its LED goes out.
The message remains on the screen.
- 3 Alarms which can be acknowledged via Alarm Reset must be acknowledged via the »Alarm Reset« key, see "Fault – Cause – Remedy", page 84.

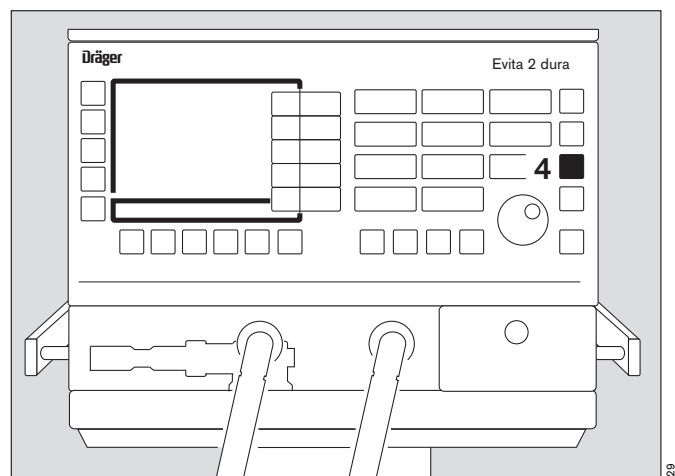


Information i

- For help with system operation
 - For help with troubleshooting
- 4 Press the »Info« key: the required information is displayed in the bottom line of the screen.

To erase the message:

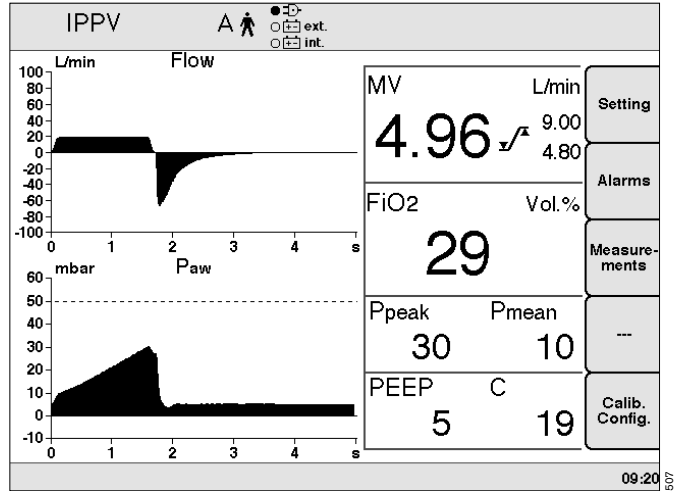
- 4 Press the »Info« key again.



Displaying Curves and Measured Values

In the main page

A set of six selectable measured values is displayed in the right-hand field, and two selectable curves in the left-hand field.



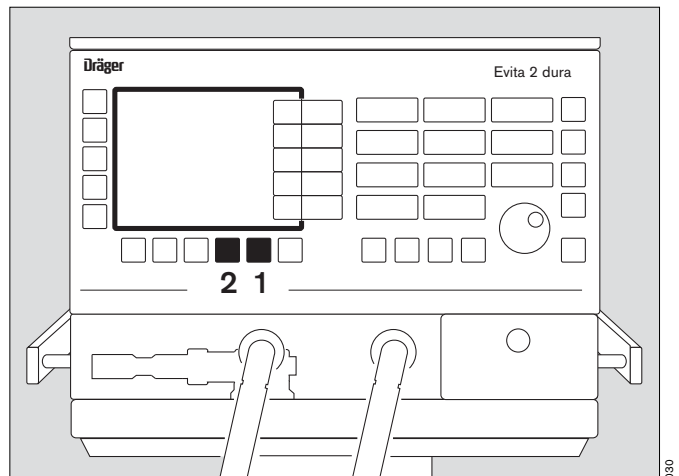
To select a second group of 6 measured values:

- 1 Press the »Values 1 2 « key.

To select another pair of curves:

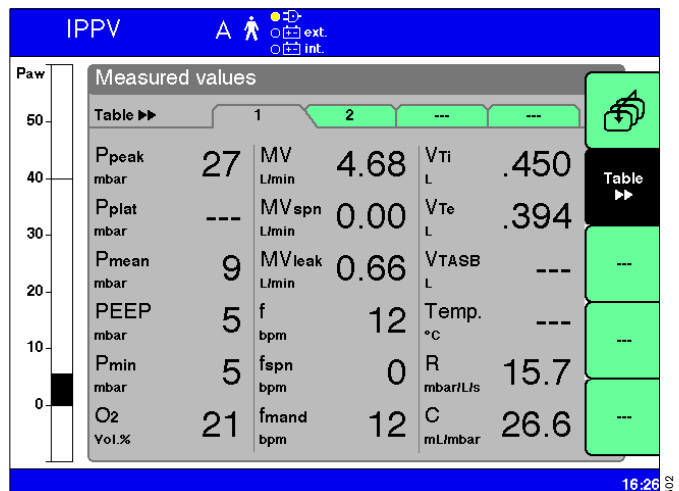
- 2 Press the »Curves « key.

Measured values and curves can be selected, see "Combine displayed measured values", page 75 and "Combine displayed curves", page 76.



In the other screen pages, these curves and measured values are not displayed. The airway pressure is therefore continuously indicated by means of an analogue vertical bar display on the left-hand side of the screen.

Example: screen page »Measures values 1«



Display measured values

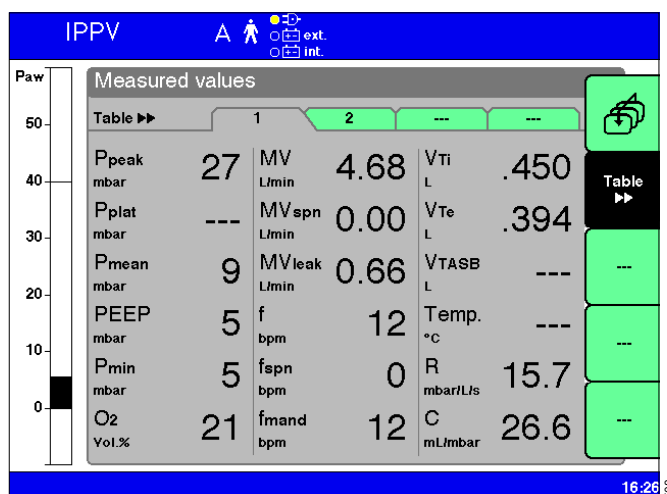
- Press menu button »Measurements«.

Display example: »Table 1«

Evita 2 dura displays the measured values and their units of measure in the form of a table. The menu »Table 1« is displayed with all standard available measured values.

The measured value MV_{leak} represents the leakage in L/min and is determined by Evita 2 dura by comparing the applied inspiratory minute volume with the measured expiratory minute volume.

The measured value MV_{leak} is used by Evita 2 dura for automatic correction of the applied tidal volume V_{Ti} and the flow and volume curves. This presupposes that leakage compensation has been activated, see page 80. For safety reasons, the measured values for the minute volume are not corrected.

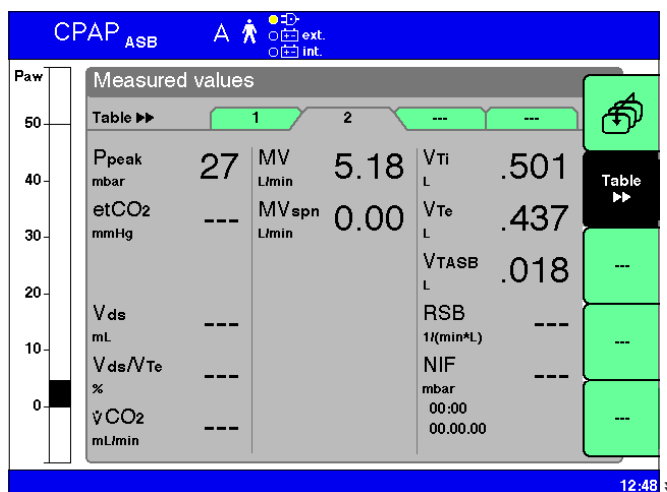


To display the measured values of add-on features (e.g. "Evita 2 dura CapnoPlus"):

- Select the »Table 2« menu with the »Table ►►« menu key.
- Functional extensions of measured values, e.g. CapnoPlus CO₂ monitoring, are displayed in further measured value tables.

The following are optionally displayed (upgrade SW4.n plus)

- VTASB Inspiratory tidal volume during an ASB stroke
- RSB Rapid Shallow Breathing*
- NIF Negative Inspiratory Force**



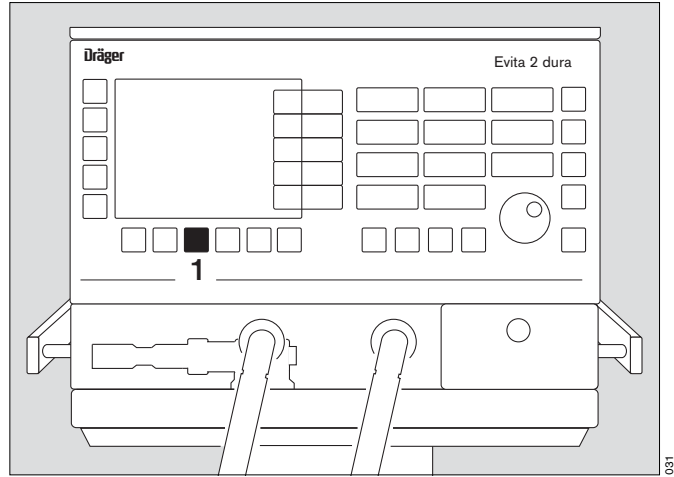
* For a detailed description of RSB, see page 126.

** For a detailed description of NIF, see page 126.
Use of NIF, see "Manual expiration", page 59.

Curve freezing

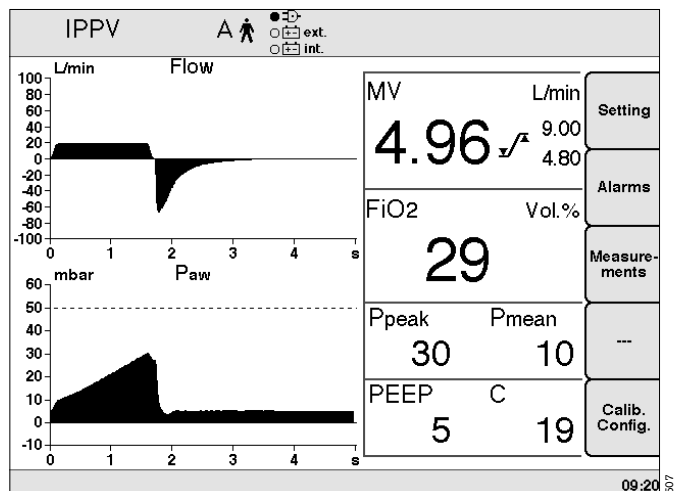
To study the curve(s) in detail:

- 1 Press the »Freeze « key.



To return to displaying new curve(s):

- 1 Press the »Freeze « key again.



Special Functions

Manual inspiration

This function may be used in all modes except CPAP spontaneous breathing without ASB pressure support. An automatic ventilation stroke can be triggered independently of the starting time and extended up to a maximum of 15 seconds.

Or:

Between two automatic ventilation strokes, a ventilation stroke can be manually started and held for a maximum 15 seconds.

The pattern of the manually started ventilation stroke depends on the ventilation mode used.

For IPPV, SIMV and MMV:

volume-controlled ventilation stroke, defined by the VT and T_{insp} settings.

For BIPAP:

pressure-controlled ventilation stroke, defined by the P_{insp} and T_{insp} settings.

For CPAP/ASB:

pressure-controlled ventilation stroke, defined by the PASB setting.

- 1 Press and hold down the »**Insp. hold**« key for as long as inspiration is required.

Either an automatic ventilation stroke that has just begun will be prolonged for as long as the key is held down, or a new ventilation stroke will be started and prolonged for as long as the key is held down – in each case for a maximum of 15 seconds.

Manual expiration hold

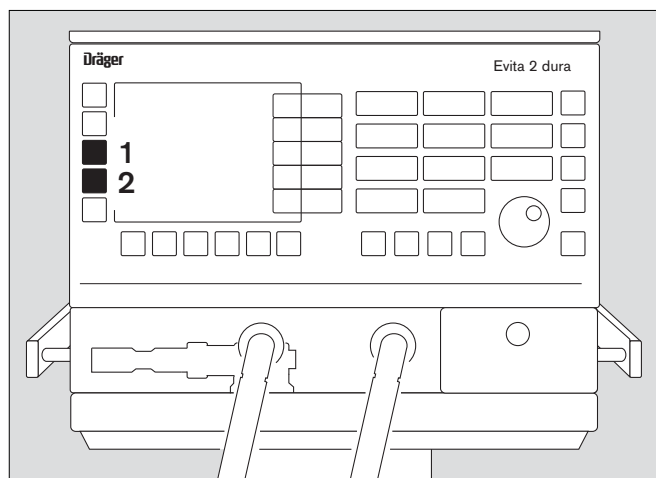
Active in all ventilation modes.

For determining the weaning value NIF*

- 2 Hold down the »**Exp. hold**« key.

The expiration phase remains effective and Evita 2 dura determines the measured NIF value as long as the key is pressed.

After 15 seconds, the system automatically interrupts the expiration phase.



* For a detailed description of NIF, refer to the Annex, page 126.

Medicament nebulisation

Inflammable substances must not be nebulised! Fire hazard due to the glowing flow sensor. Nebulants containing flammable substances such as alcohol must NOT be used, any contact of a flammable substance with the hot wire flow sensor could cause a fire.

During adult ventilation

Applicable in every ventilation mode.

Evita 2 dura applies the medicament aerosol in synchronisation with the inspiratory flow phase and maintains the minute volume constant.

Depending on the set O₂ concentration, the ventilator supplies the medicament nebuliser with medical air, pure oxygen or a mixture of medical air and oxygen. Deviations in O₂ concentration are therefore kept to a minimum.

In extreme cases (with a minimum inspiration flow of 15 L/min), the deviations can be up to $\pm 4\%$ by volume*. To avoid greater deviations, medicament nebulisation is automatically switched off with inspiration flows of less than 15 L/min.

During paediatric ventilation

Medicament nebulisation is possible in the pressure-controlled paediatric ventilation modes.

In volume-controlled ventilation modes, medicament nebulisation is only possible with AutoFlow[®] (optional extra).

Contrary to the case during adult ventilation, the medicament is nebulised continuously in paediatric mode. However, the aerosol generated during expiration does not enter the lungs. Nebulising while in paediatric mode is continuous, please note that nebulised medication will not enter the lungs during the expiratory phase of the breath. Nebulising in the adult mode is synchronized with the inspiratory phase only, and is not active during the expiratory phase.

Depending on the set O₂ concentration, the medicament nebuliser is supplied by the ventilator with medical air, oxygen or a mixture of medical air and oxygen. Deviations in O₂ concentration are therefore kept to a minimum.

We recommend that you do not use the medicament nebuliser at breathing rates of less than 12 bpm!

For breathing rates above 12 bpm, please refer to the graph on page 127 of these Instructions for Use.

The maximum possible deviations in O₂ concentration are $\pm 4\%$ by volume.

For breathing rates of less than 12 bpm, the deviations in O₂ concentration may be much greater. These deviations cannot be detected by the device's internal O₂ concentration monitor.

* For a detailed description of the inspiratory O₂ concentration during (medicament) nebulisation, see page 127.

The minute and tidal volumes displayed may be considerably higher or lower than those actually set to be delivered to the patient due to varying tolerances in flow during nebulising. The measured values prior to nebulisation must be taken into account if necessary.

In the event of a discrepancy between VT and MV values, the ventilation pressure can be used for an accurate assessment of ventilation. VT and MV values can be compared by comparing the difference between PEEP and plateau pressure before and during nebulisation.

The medicament nebuliser is automatically switched off after 30 minutes. After administration of the aerosol, the flow sensor is automatically cleaned and calibrated in order to prevent malfunctions in flow measurement.

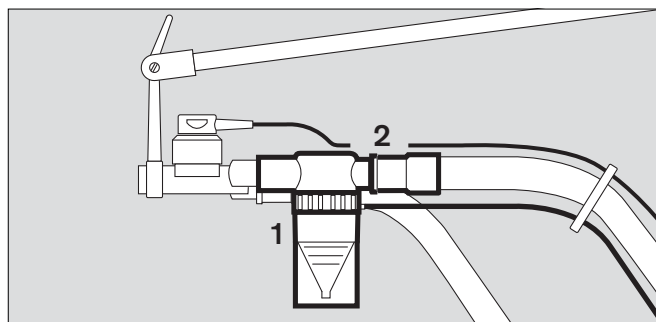
Use only medicament nebuliser 84 12 935 (white middle section).

Prepare the medicament nebuliser as specified in the specific Instructions for Use.

If other pneumatic medicament nebulisers are used, major deviations in tidal volume and inspiratory O₂ concentration may be caused!

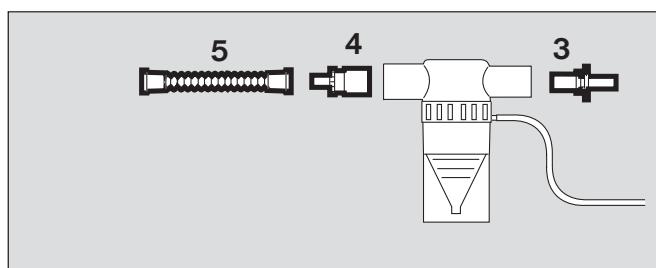
For use during adult ventilation

- 1 Connect the nebuliser to the inspiratory side (temperature sensor side) of the Y-piece.
- 2 Connect the inspiration hose to the medicament nebuliser.
- Place the medicament nebuliser in the vertical position.
- Using hose clips, route the nebuliser hose back to the ventilator along the expiratory hose.

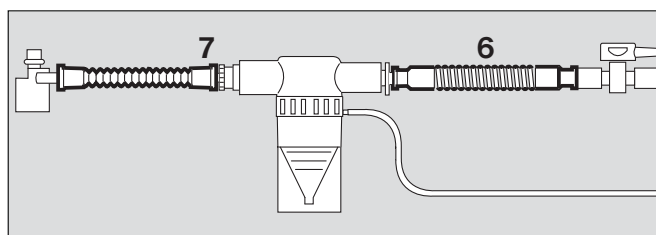


For use during paediatric ventilation

- 3 Insert the catheter connector (ISO cone Ø 15/Ø 11) into the inlet of the medicament nebuliser.
- 4 Insert the adapter (ISO cone Ø 22/Ø 11) into the outlet.
- 5 Fit the corrugated hose (0.13 m long) on to the outlet adapter.



- 6 Remove the corrugated hose of the hose set from the inspiratory adapter of the Y-piece and connect it to the inlet adapter of the medicament nebuliser.
- 7 Connect the free end of the corrugated hose on the nebuliser outlet to the inspiratory adapter of the Y-piece.



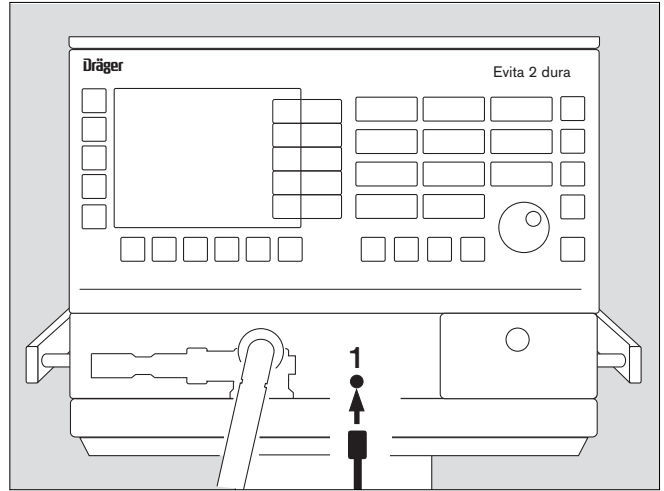
- 1 Connect the nebuliser hose to the port on the front panel of the Evita 2 dura.
- Fill the medicament nebuliser in accordance with its specific Instructions for Use.


Warning: the effect of aerosols on sensors, filters and heat and moisture exchangers (HME) must be taken into account!

The measuring function of the flow sensor may be impaired. The flow resistance of filters is liable to increase and may impair ventilation.


Do not place a microbial filter on the nebuliser outlet during nebulisation!

During medicament nebulisation, do not use a heat and moisture exchanger (HME) at the Y-piece. Risk of increased breathing resistance!



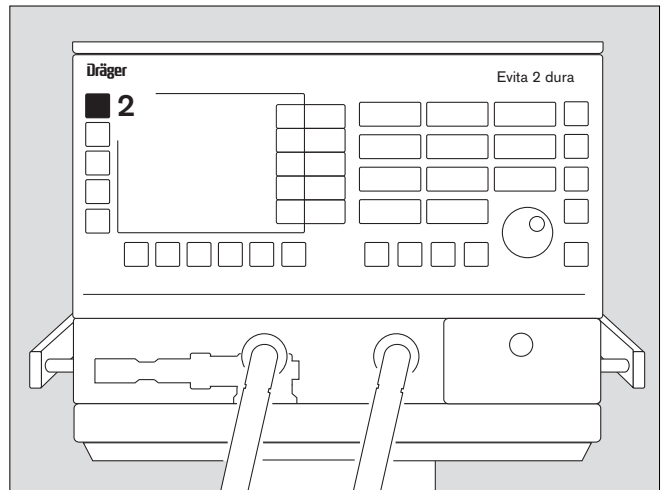
- 2 Hold down key »  « until the yellow LED lights up.
- Advisory message on the screen:
»Nebuliser on!«
The nebuliser remains in operation for 30 minutes.

If you wish to interrupt medicament nebulisation before it is complete:

- 2 Press »  « again. The yellow LED goes out, the nebuliser will be switched off.

The flow sensor is subsequently cleaned and calibrated automatically.

- Remove residual medicament. Strictly follow the Instructions for Use of the medicament nebuliser.



Oxygen enrichment for bronchial suction

To avoid any risk of hypoxia during bronchial suction, Evita 2 dura offers a programme for oxygen enrichment during the removal of secretions.

After the programme is started, Evita 2 dura ventilates the patient in the selected ventilation mode for an initial oxygen enrichment phase of 180 seconds. In adult mode, the ventilator supplies 100 % oxygen by volume, and in paediatric mode it delivers the set O₂ concentration* plus 25 % (for example: setting = 60 % by vol.; administered = 75 % by vol.)

When the ventilator is disconnected for suction, Evita 2 dura interrupts the ventilation. During the suction time, the audible alarms are suppressed, so that the suction routine is not disturbed.

After suction and automatically recognised reconnection, Evita 2 dura delivers an increased O₂ concentration* for the final oxygen enrichment phase of 120 seconds. In adult mode, the O₂ concentration is 100 % by volume. In paediatric mode, the enriched concentration is 25 % higher than the set concentration*.

During suction and for 2 minutes afterwards, the lower alarm limit for the minute volume is switched off.

Before suction

- 1 Hold down the »O₂ ↑ Suction« key until the yellow LED comes on.

Evita 2 dura ventilates the patient in the set ventilation mode with increased O₂ concentration: 100 % O₂ by volume in adult mode, and a 25 % higher O₂ concentration than the set value in paediatric mode.

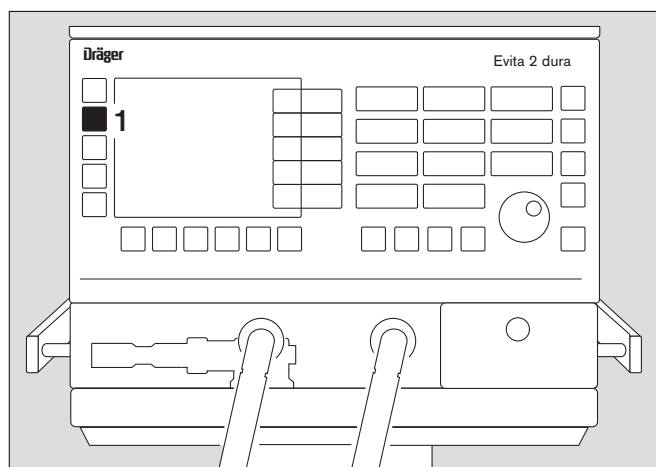
If no PEEP greater than 4 mbar is set, a PEEP of 4 mbar will be automatically activated. This PEEP enables the Evita 2 dura to detect subsequent disconnection.

The other ventilation parameters remain unchanged.

- Display in the help line at the bottom edge of the screen:
»O₂ enrichment 180 s«

The remaining time is counted down continuously. This initial oxygen enrichment lasts for a maximum of 180 seconds. During this time, Evita 2 dura waits for a disconnection for suction.

If there is no disconnection after expiry of the 180 seconds, the oxygen enrichment programme is automatically terminated.



* For a detailed description of the inspiratory O₂ concentration during medicament nebulisation, please refer to the Appendix page 127.

After disconnection for suction

Evita 2 dura delivers a minimal flow for the duration of disconnection in order to detect the end of the disconnection phase automatically.

In the help line at the bottom of the screen, the amount of time available for suction is continuously counted down (example):
»Execute suction and reconnect 120 s«

If suction is ended and ventilation reconnected within the displayed time, Evita 2 dura starts the final O₂ enrichment phase.

Automatic interruption of oxygen enrichment

If there is still no reconnection after 120 seconds, the oxygen enrichment programme is reinitiated. All alarms are immediately reactivated. Evita 2 dura immediately continues ventilating in the set ventilation mode.

After reconnection

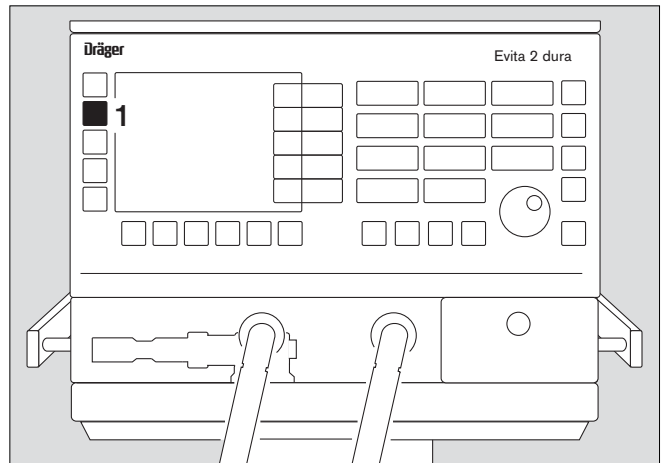
Evita 2 dura continues ventilating in the set ventilation mode, except that for final oxygen enrichment the increased oxygen concentration of 100 % by volume for adults and 25 % above the set concentration for paediatric ventilation will continue to be delivered for 120 seconds.

- Message in the help line at the bottom of the screen:
»Final O₂ enrichment 120 s«

The time remaining is counted down continuously.

To interrupt oxygen enrichment

- 1 Press the »O₂ ↑ Suction« key again.





Selecting Standby Mode

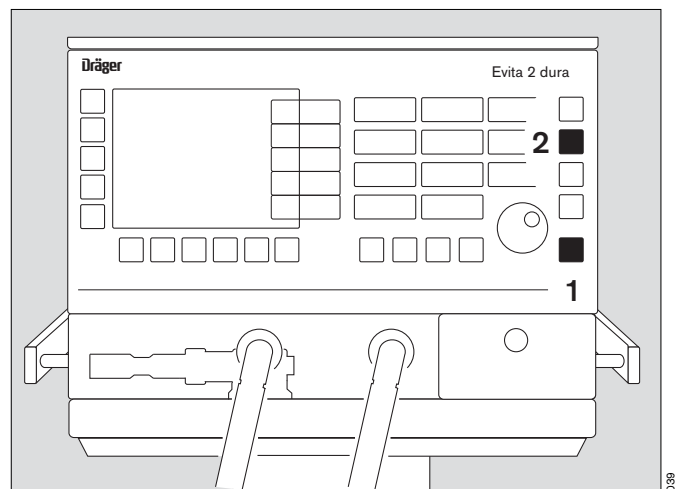
No ventilation takes place in standby mode!

- to select the patient mode
- to perform the device check
- to maintain Evita 2 dura ready for operation
- to preset ventilation parameters and alarm limits.


Switching to Standby

- 1 Hold down the »« key for about 3 seconds. The Standby alarm tone is sounded.
- 2 The »Alarm Reset« key can be used to switch off the Standby alarm tone.

The standby alarm tone can no longer be muted with the »« key.

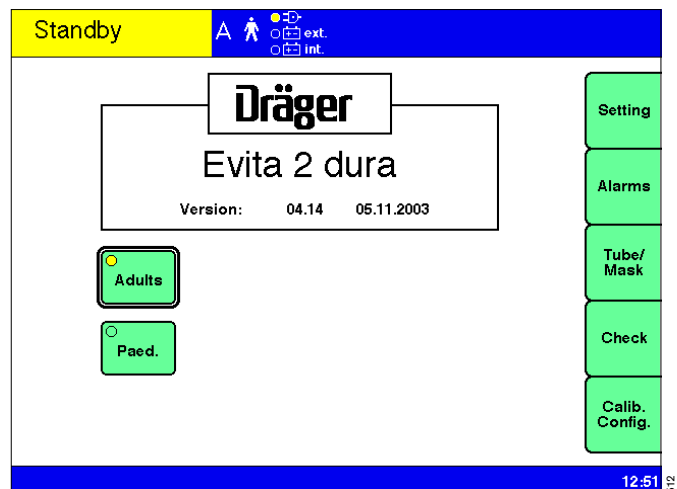


Terminating standby mode

- to continue ventilation.
- 1 Press »« key. The LED goes out, and ventilation commences.

If the patient mode is changed while on standby, Evita 2 dura will determine new starting values for ventilation, see page 78.

Display (example):



Calibration

The saved calibration/zeroing values remain stored even when the machine is switched off.

The pressure sensors for measuring the airway pressure are calibrated automatically.

The flow sensor and O₂ sensor are automatically calibrated once per day.

Calibration of the flow sensor can be performed at any time, even during ventilation.

Calibration of the O₂ sensor can be performed at any time, even during ventilation. The applied O₂ concentration is not affected by calibration.

The calibration of the CO₂ sensor (optional) can be checked during ventilation.

Calibrating the O₂ sensor manually

- Before operation, during the device check.
- After replacing the O₂ sensor (wait for the 15-minute warm-up time of the O₂ sensor).
- If the measured value and set value deviate from each other by more than 2 Vol.%.

The O₂ sensor can be calibrated during ventilation.

Start calibration:

- Press the »Calib./Config.« menu key.
- Select the »O₂« menu key = turn the rotary knob.
- Start »O₂« calibration = press the rotary knob.

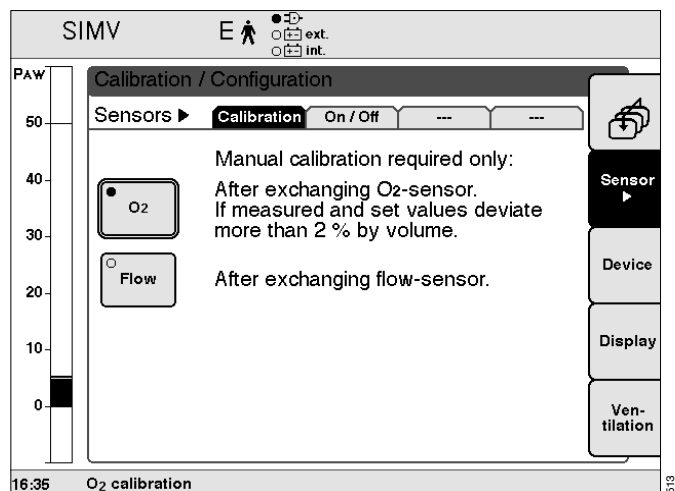
Display (example):

Message in the help line at the bottom of the screen:

»O₂ calibration«

After calibration is complete, the following message appears in the help line:

»Calibration ok«



Calibrating the flow sensor manually

- Before operation, during the device check.
- After replacing the flow sensor.

The flow sensor is automatically cleaned before each calibration.

After using the medicament nebuliser, the flow sensor is automatically cleaned and calibrated.

To start calibration:

- Avoid flammable gases (e.g. alcohol vapours after disinfection).
- Flow sensors which have been disinfected in ethanol must be left to dry in air for at least 30 minutes.
- Press the »Calib./Config.« key.
- Select the »Flow« screen key = turn the rotary knob.
- Start calibration = press the rotary knob.

Display (example):

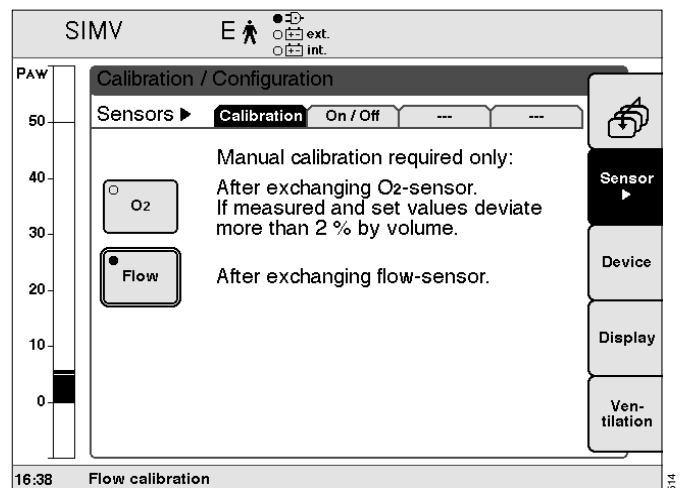
Evita 2 dura uses the next inspiration phase for the calibration. Short inspiration times are prolonged to about 1 second.

Message in the help line at the bottom of the screen:

»Flow calibration«

After calibration is complete, the following message appears in the help line:

»Calibration ok«



External flow source

When a constant external flow of up to 12 L/min is applied (e.g. during medicament nebulisation with separate gas supply and not from Evita 2 dura or during separate tracheal gas insufflation TGI), this flow can be determined by Evita 2 dura and the tolerance for the flow sensor monitoring parameters increased in order to prevent generation of the alarms "Flow measurement fault" and "Neo. flow measurement fault" (NeoFlow option) during these applications.

The original measurement of the expiratory volume is continued: During an expiratory flow, Evita 2 dura measures a correspondingly higher value for V_{Te} and MV.

The displayed V_{Ti} value is too low. During volume-controlled ventilation, the actual tidal volume applied to the patient is higher than that set. It is therefore advisable to use pressure-controlled ventilation in combination with an external flow.

To avoid false alarms and ensure adequate monitoring:

- Adjust both alarm limits for MV in line with the actual value.
- Use additional monitoring, e.g. SpO₂, if necessary.

To determine the external flow:

- Press menu button »Calib./Config.«.
- Select the menu »External Flow« via the menu button »Sensor ▶▶«.
- Select screen key »Measure« = turn rotary knob, confirm = press rotary knob.
- Yellow LED lights up in the »Measure« key.

The external flow is calculated by Evita 2 dura.

Display during calculation:

»Determining external flow«

Once the external flow has been determined, it is displayed by Evita 2 dura together with the time and date.

The following message is simultaneously displayed by Evita 2 dura:

»Confirm value via «

- Confirm = press rotary knob.

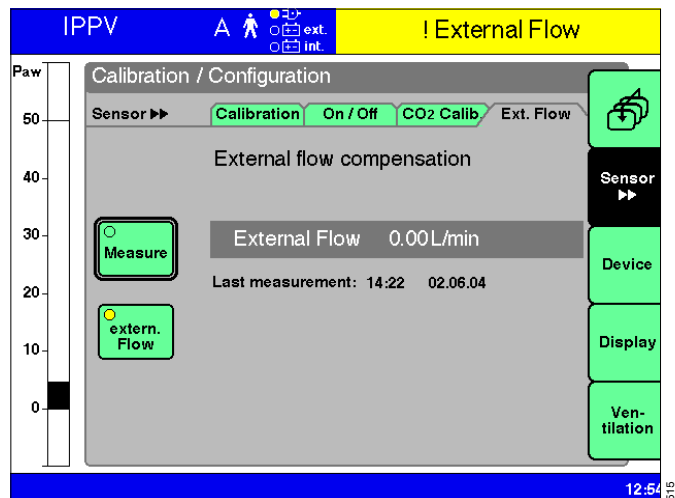
Determination of the external flow is aborted by Evita 2 dura if it exceeds 12 L/min or if flow measurement by Evita 2 dura is faulty.

When the external flow has been determined successfully, it is taken into account in the monitoring of the flow sensor: the yellow LED in the key »extern. Flow« lights up.

The advisory message

»! External flow«

is displayed as long as the external flow is taken into account by Evita 2 dura.



When an external flow is not applied:

- Switch off:
select key »**extern. Flow**« = turn rotary knob,
confirm = press rotary knob.

Once the external flow has been measured by Evita 2 dura,
its inclusion can be reactivated at any time:

- Select screen key »**extern. Flow**« = turn rotary knob,
confirm = press rotary knob.

If the external flow changes:

- Press key »**Measure**« so that the external flow can be
redetermined by Evita 2 dura.

Switching off the monitor functions

E.g. if a spent sensor cannot be immediately replaced.

**An adequate external monitoring function must
immediately be ensured!**

Example: Switching off Flow Monitoring

- Press the »**Calib./Config.**« menu key.
- Select »**Sensor On/Off**« with the »**Sensor ▶▶**« menu key.

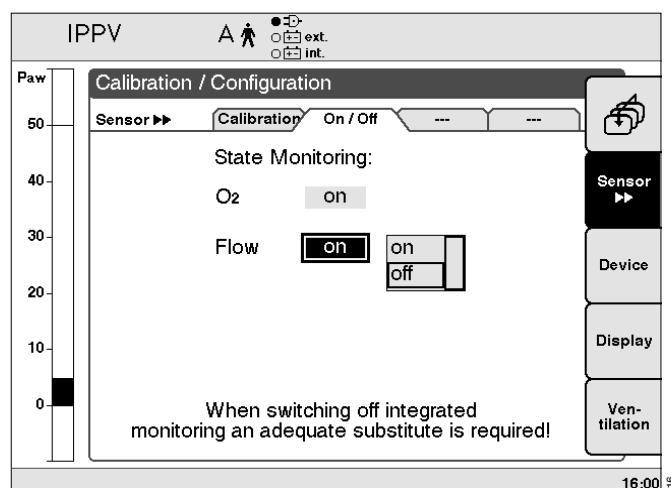
Example display:

- Select the »**Flow on**« screen line = turn the rotary knob.
Confirm = press the rotary knob.
- In the selection menu, select »**off**« = turn the rotary knob.
Confirm = press the rotary knob.

The corresponding measured values disappear.
The alarm function is deactivated.

To switch the monitor function back on after replacing the
sensor:

- Select »**Flow off**« screen line = turn the rotary knob.
Confirm = press the rotary knob.
- In the selection menu, select »**on**« = turn the rotary knob.
Confirm = press the rotary knob.



Configuration

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Configuration

System Settings

Adjusting the volume of the audible alarm

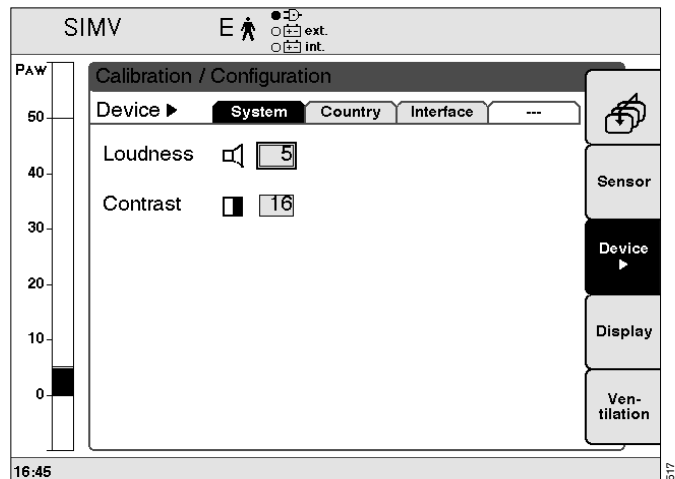
- Press the »Calib./Config.« menu key.
- Press the »Device ►►« menu key. The »System« menu appears.

Display (example):

- Select the »Loudness« screen field = turn the rotary knob.
Activate = press the rotary knob.
- Set the desired volume = turn the rotary knob.
Confirm = press the rotary knob.

After the setting has been confirmed, the alarm tone is sounded once to enable you to judge the volume.

Set the volume of the acoustic alarm to a sufficiently high level so that the alarm can be heard!



Setting the contrast

Not possible with all types of monitor

- Press the »Calib./Config.« menu key.
- Press the »Device ►►« menu key. The »System« menu is displayed.
- Select the »Contrast« field on the screen = turn the rotary knob.
Activate = press the rotary knob.
- Setting the contrast = turn the rotary knob,
confirm = press the rotary knob.

The set contrast will now be activated.

Country-specific settings

Selecting the language

Evita 2 dura is supplied in the language of the customer's country.

The following languages can be selected:

- English
- French
- Italian
- Spanish
- Dutch
- Swedish
- American English
- Japanese
- Greek
- Russian
- Portuguese
- Arabic
- Chinese
- Turkish

- Press the »Calib./Config.« menu key.
- Press the »Device ►« menu key.
- With the »Device ►« menu key, select the »Country« menu.

Display (example):

- Select the »Language« screen field = turn the rotary knob.
Confirm = press the rotary knob.
- Select language = turn the rotary knob.
Confirm = press the rotary knob.

Setting the date and time

- Press the »Calib./Config.« menu key.
- Press the »Device ►« menu key.
- With the »Device ►« menu key, select the »Country« menu.
- Select the »Day« screen field = turn the rotary knob.
Confirm = press the rotary knob.
- Set the date = turn the rotary knob.
Confirm = press the rotary knob.

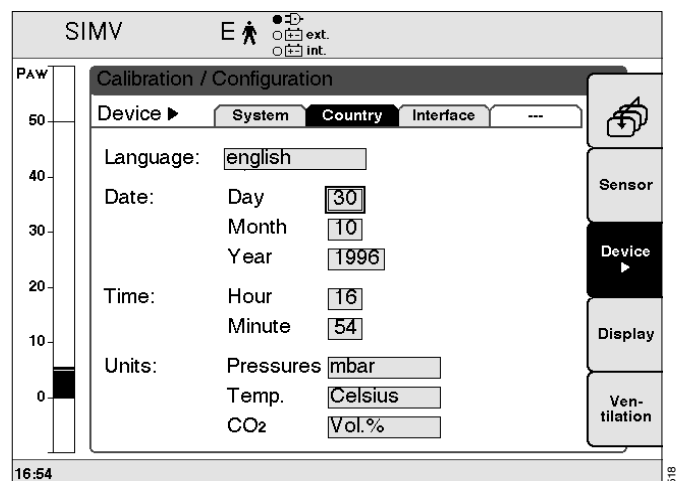
Set the month, year, hour and minute in the same way.

Selecting measuring units

- Press the »Calib./Config.« menu key.
- Press the »Device ►« menu key.
- With the »Device ►« menu key, select the »Country« menu.

Under units:

- Select the »Pressures« screen field = turn the rotary knob.
Confirm = press the rotary knob.
- Set »Temp.« and »CO2« (option) in the same way.



Interface

Selecting the interface

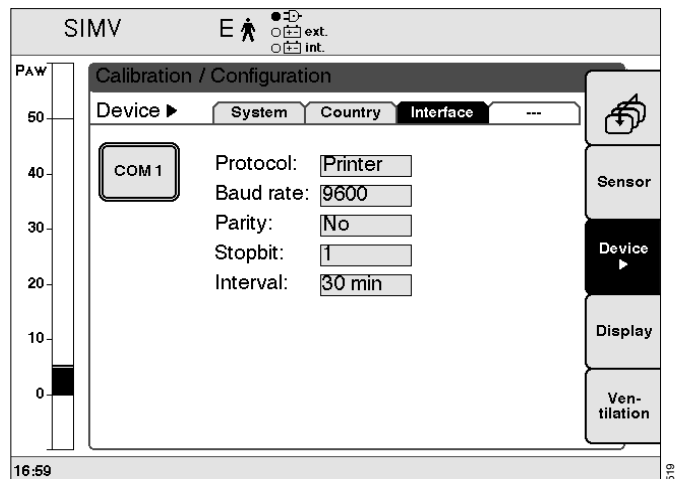
Evita 2 dura offers the following interface protocols:

- Printer (HP Deskjet 500 Series 500 and compatible printers with serial interface)
- MEDIBUS (Dräger communications protocol for medical appliances)
- LUST (List-controlled universal interface driver program, compatible with the Evita RS 232 interface as from software version 7.n)

Other equipment, e.g. printers, may only be connected to the COM ports if Evita 2 dura is connected to the mains power supply via a mains power cable or if it has been earthed to the earth connection on the back of the unit.

Electric power may pose a hazard in all other cases.

- Press the »Calib./Config.« menu key.
- Press the »Device ▶« menu key.
- With the »Device ▶« menu key, select the »Interface« menu.
- Select the screen key corresponding to the required interface, »COM1«, »COM2«, »COM3« and »Analog« = turn the rotary knob.
Confirm = press the rotary knob. (COM2, COM3 and Analog are optional).
- Select the desired interface protocol in the "Protocol" screen field = turn the rotary knob.
Confirm = press the rotary knob.
- Select the screen field corresponding to the desired interface parameter = turn the rotary knob.
Confirm = press the rotary knob.
- Set the desired value = turn the rotary knob.
Confirm = press the rotary knob.



Adapting the interface protocols:

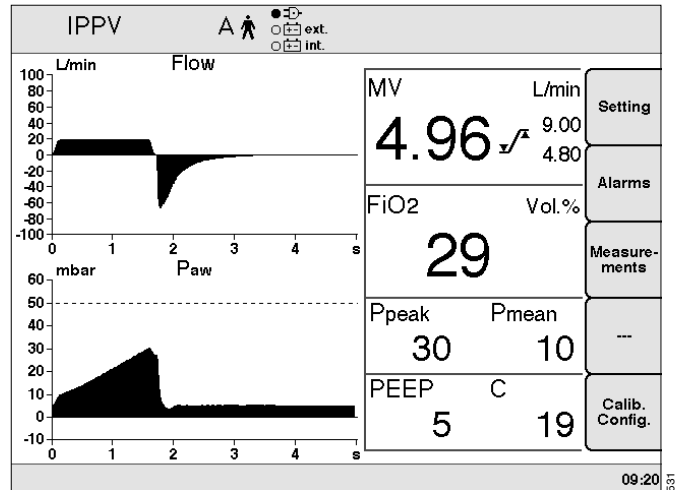
- See the Instructions for Use of the device you want to connect.
For the printer protocol:
Baud rate
Set printer interval as required
- For the MEDIBUS protocol:
Baud rate
Parity check bits
Number of stop bits
- For the LUST protocol:
Baud rate

Screen

Selecting the displayed combination of measured values

Evita 2 dura displays a group of 6 selectable measured values in the right-hand field of the main page.

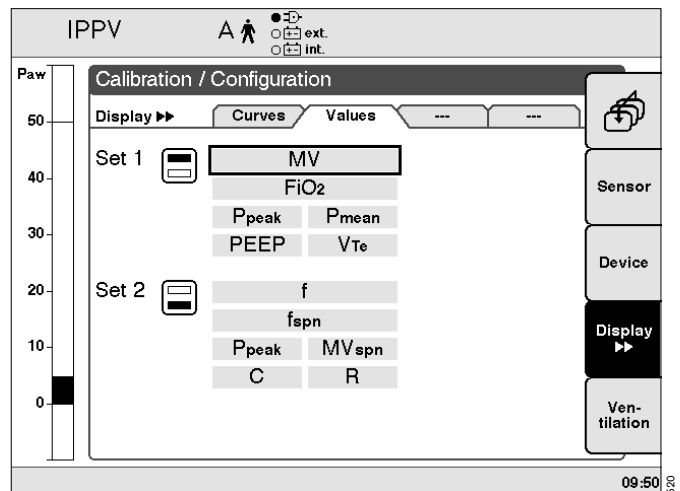
An alternative second group can be displayed by pressing the »Values 1 □ 2 □ «.



These two groups can be composed in the configuration page:

- Press the »Calib./Config.« menu key.
- Press the »Display ►►« menu key.
- With the »Display ►►« menu key, select the »Values« menu.

Display (example):



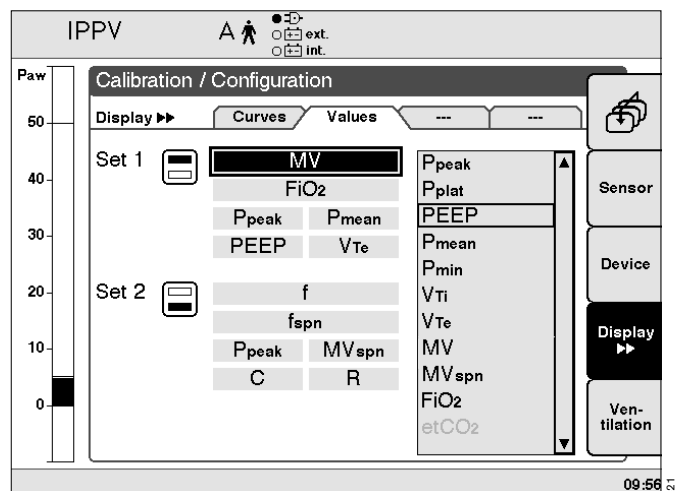
To replace one displayed measured value by another:

- Select the relevant screen field = turn the rotary knob.
Activate = press the rotary knob.

The selection list with all available measured values is then displayed on the right of the screen.


Display (example): Replacing MV

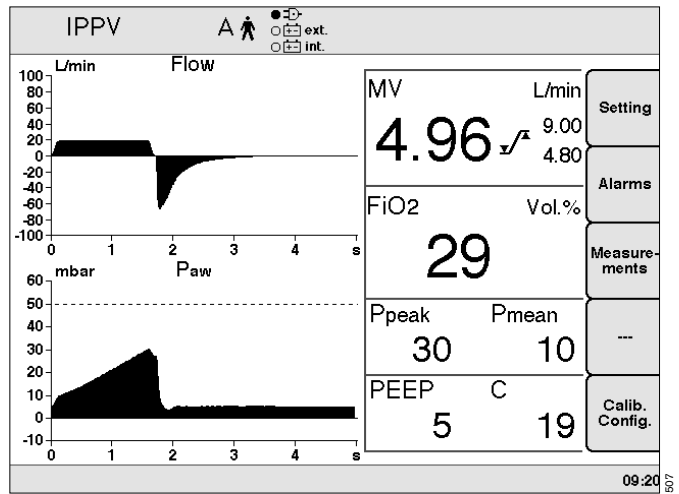
- Select the other measured value, e.g. »PEEP« = turn the rotary knob.
Confirm = press the rotary knob.




Selecting the displayed curves

Evita 2 dura shows two curves in the left-hand field of the main page.

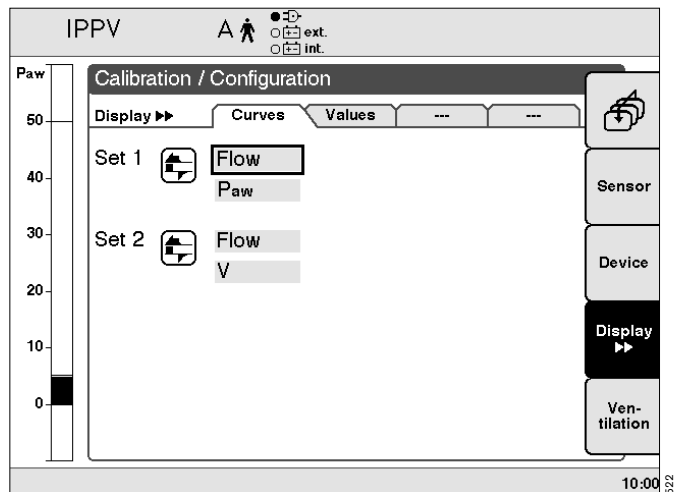
A different pair of curves can be selected by pressing the »Curves « key.



The curve pairs can be combined as required.

- Press the »Calib./Config.« menu key.
- Press the »Display « menu key. The »Curves« menu appears.

Display (example):



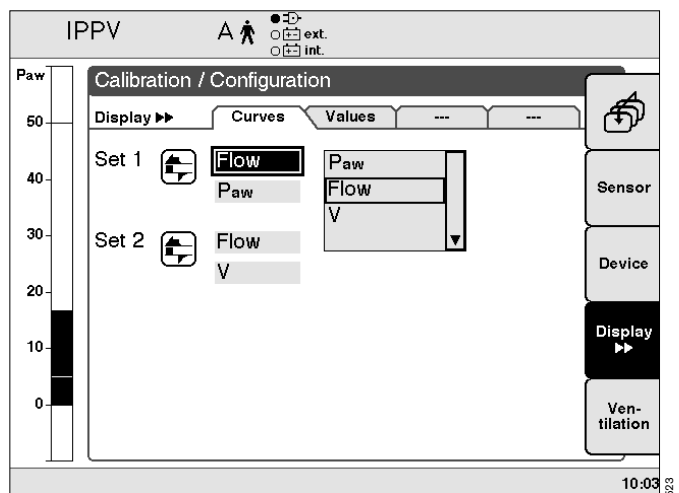
One displayed curve can be replaced by another:

- Select the relevant field = turn the rotary knob.
Activate = press the rotary knob.

The list of all available curves to choose from is displayed on the right-hand side of the screen.

Display example: Replacing Paw

- Select the other curve (»Flow«) = turn the rotary knob.
Confirm = press the rotary knob.

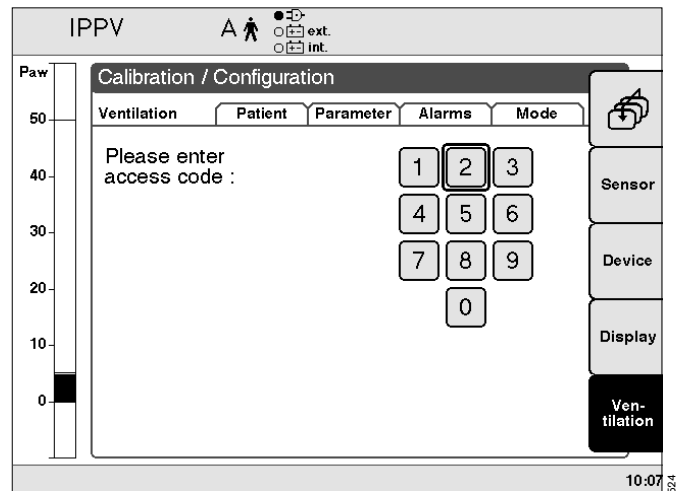


Ventilation Defaults

- For defining the patient-specific start-up parameters »f« and »VT«
- Ventilation parameters active on starting up the device
- Alarm limits active on starting up the device
- Ventilation mode

The »Ventilation« menu for the default settings active on starting ventilation can only be accessed after entering the code number 3032.

This code-protection is designed to prevent accidental changes to the configuration.



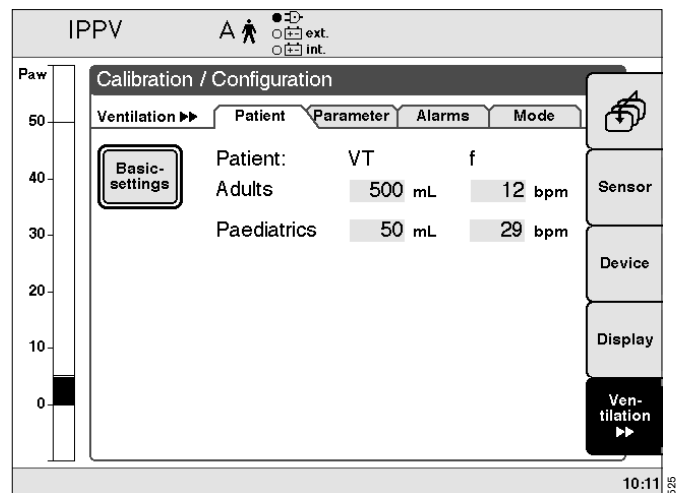
Patient-specific defaults

for Adult or Paediatric mode

- To set the values of the »f« and »VT« parameters active on starting up the device.
- Press the »Calib./Config.« menu key.
- Press the »Ventilation ►►« menu key.
- Enter code number »3032«.
- The »Patient« menu is displayed.

Display (example):

- Select the »VT« screen field = turn the rotary knob.
Confirm = press the rotary knob.
- Set the desired value = turn the rotary knob.
Confirm = press the rotary knob.



Patient-specific defaults VT, f

Patient mode	Factory setting		Hospital-specific setting	
	Tidal volume VT mL	Ventilation frequency f 1/min	Tidal volume VT mL	Ventilation frequency f 1/min
Paediatric	50	29		
Adult	500	12		

The hospital-specific defaults for your hospital can be entered in the table.

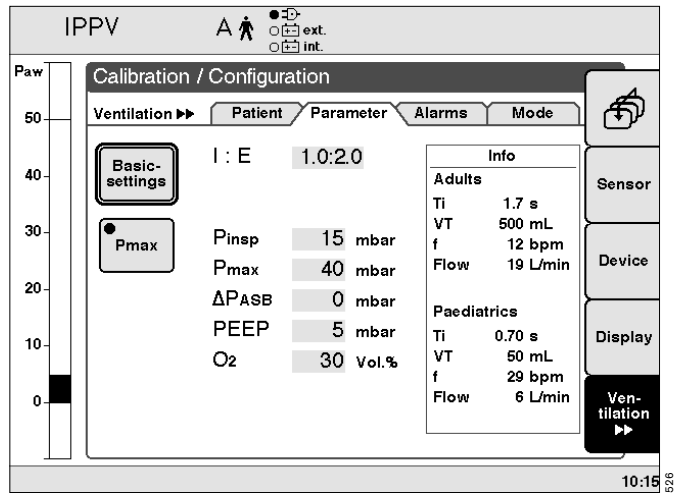
Default ventilation parameter values

– These defaults specify the ventilation parameters and alarm limits that are not patient-specific and are activated on starting up the device.

- Press the »Calib./Config.« menu key.
- Press the »Ventilation ►►« menu key.
- Enter code number »3032«.
- Select the »Parameter« menu with the »Ventilation ►►« menu key.

Display (example):

- Select the Pmax desired screen field = turn the rotary knob.
Confirm = press the rotary knob.
- Set value = turn the rotary knob.
Confirm = press the rotary knob.



Default ventilation parameters

	I:E	P _{insp} mbar	PASB mbar	PEEP mbar	Ramp s	Trigger L/min	O ₂ Vol.%
Factory setting	1:2	15	0	5	0.2	5	30
Hospital-specific setting							

The hospital-specific defaults for your hospital can be entered in the table.

To restore the factory-set defaults:

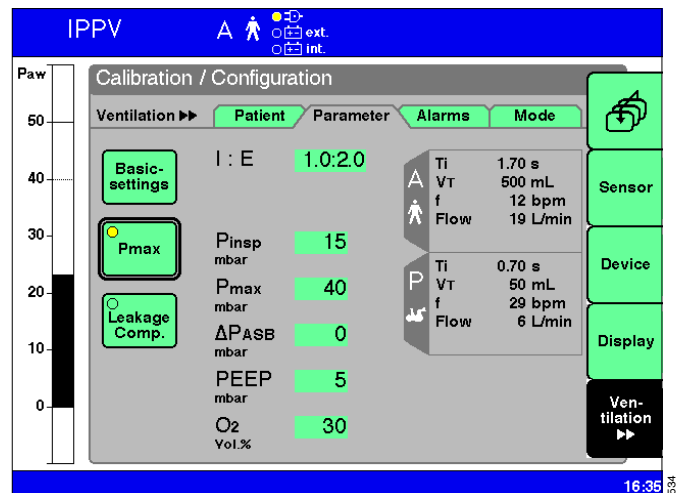
- Select the »Basic settings« screen field = turn the rotary knob.
Confirm = press the rotary knob.

Activating/deactivating pressure limit Pmax

- This parameter defines the pressure limit for pressure-limited ventilation in the IPPV, SIMV and MMV ventilation modes.
- Press the »Calib./Config.« menu key.
- Press the »Ventilation ►►« menu key.
- Enter code number »3032«.
- With the »Ventilation ►►« menu key, select the »Parameter« menu.

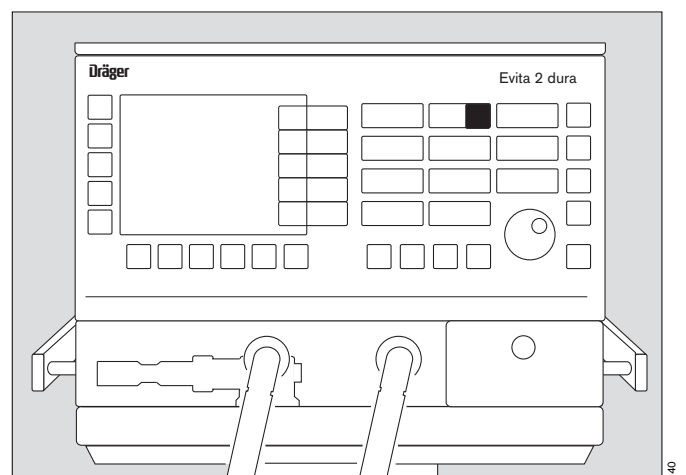
Display (example):

- Select the »Pmax« screen key = turn the rotary knob. The "LED" in the screen key »Pmax« lights up yellow. Pmax is activated.
- Deactivate Pmax = press rotary knob, the "LED" in the screen key »Pmax« lights up green.



Set the value for Pmax:

- Press the »P_{insp}« parameter key.
- Set the desired value = turn the rotary knob.
- Confirm = press the rotary knob.



Apnoea ventilation on/off

To determine whether apnoea ventilation is automatically ready for use when starting.

- Press menu button »Calib./Config.«.
- Press menu button »Ventilation ►►«.
- Enter code number »3032«.
- Select the menu »Mode« via the menu button »Ventilation ►►«.
- Select screen key »Apnoea vent.« = turn rotary knob.
- Switch on apnoea ventilation = press rotary knob, the "LED" in the screen key »Apnoea vent.« lights up yellow.
- Switch off apnoea ventilation = press rotary knob, the "LED" in the screen key »Apnoea vent.« lights up green.

Leakage compensation* on/off

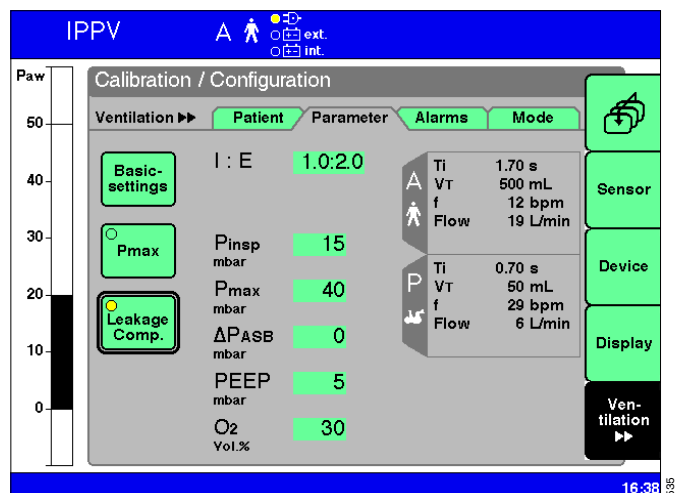
Automatic leakage compensation allows the unit to compensate leakages up to 100 % of the set tidal volume in all volume-controlled ventilation modes.

The setting for »Leakage compensation on/off« is saved and remains effective when the unit is restarted.

- Press menu button »Calib./Config.«,
- Press menu button »Ventilation ►►«.
- Enter the numerical code »3032«.
- Select the menu »Parameter« via the menu button »Ventilation ►►«.

Display:

- Select screen key »Leakage comp.«,
- Switch on »Leakage comp.« = press rotary knob, the "LED" in the screen key »Leakage comp.« lights up yellow.
- Switch off »Leakage comp.« = press rotary knob, the "LED" in the screen key »Leakage comp.« lights up green.



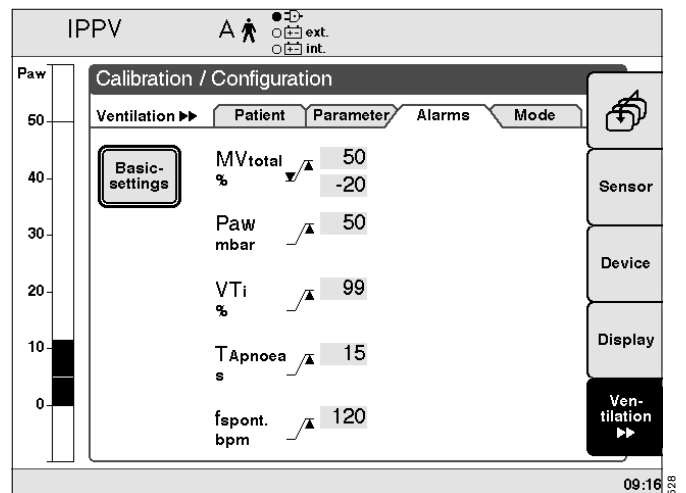
* For a detailed description of leakage compensation, refer to page 124 in the Annex.

Set initial values for alarm limits

- Press the »Calib./Config.« menu key.
- Press the »Ventilation ►►« menu key.
- Enter code number »3032«.
- With the »Ventilation ►►« menu key, select the »Alarms« menu.

Display (example):

- Select the screen field of the desired alarm limit = turn the rotary knob.
Confirm = press the rotary knob.
- Change the value = turn the rotary knob.
Confirm = press the rotary knob.



Default alarm limits

Alarm limit	Factory-set defaults	Hospital-set defaults
MVtotal low [L/min]	(VT x f) -20 %	
MVtotal high [L/min]	(VT x f) +50 %	
Paw high [mbar]	50	
VTi high [L]	VTi +100 %	
TApnoea high [s]	15	
fspont. high [1/min]	50	

The selected hospital-specific defaults can be entered in the table.

The lower alarm limit does not have to be set for the airway pressure Paw, because it is automatically coupled with the PEEP setting.

The alarm limits do not have to be set for the O₂ concentration because they are automatically coupled with the O₂ concentration setting.

Lower alarm limits:

for settings up to 60 Vol.% O₂: set value -4 Vol.% O₂
for settings from 60 to 100 Vol.% O₂: set value -6 Vol.% O₂

Upper alarm limits:

for settings up to 60 Vol.% O₂: set value +4 Vol.% O₂
for settings from 60 to 100 Vol.% O₂: set value +6 Vol.% O₂

To restore the factory-set defaults:

- Select the »Basic settings« screen field = turn the rotary knob.
Confirm = press the rotary knob.

Default ventilation mode

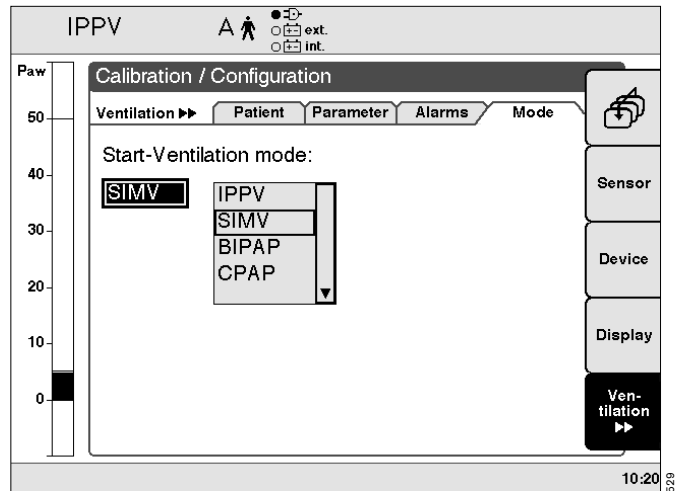
After switching on, Evita 2 dura starts up in the ventilation mode displayed when this screen key is selected. The factory-set default is IPPV.

If you require a different start-up ventilation mode:

- Press the »Calib./Config.« menu key.
- Press the »Ventilation ►►« menu key.
- Enter code number »3032«.
- With the »Ventilation ►►« menu key, select the »Mode« menu.

Display example: SIMV

- Select the screen field for the ventilation mode = turn the rotary knob.
Confirm = press the rotary knob.
The list of all available ventilation modes to choose from is then displayed on the righthand side of the screen.
- Select other ventilation mode = turn rotary knob, confirm = press rotary knob.



Fault – Cause – Remedy

Fault – Cause – Remedy 84

Fault – Cause – Remedy

Alarm messages in the alarm display field are displayed in hierarchical order.

If, for example, two faults are detected at the same time, the more critical of the two is displayed.

The priority for alarm messages is marked by exclamation marks:

Warning = Message with top priority !!!



Caution = Message with medium priority !!

Advisory = Message with low priority !

In the table below, the messages are listed in alphabetical order. The table should help you identify the cause of any alarm, and to ensure rapid remedy of the problem.

Message	Cause	Remedy
!!! Air supply down	Air supply pressure too low.	Make sure pressure is greater than 3 bar.
! Air supply down	Air supply pressure too low. Air supply pressure not required when FiO ₂ = 100 Vol.%. <hr/>	Make sure pressure is greater than 3 bar.
!! Air supply pressure high	Air supply pressure too high.	Ensure pressure is less than 6 bar.
! Air supply pressure high	Air supply pressure too high. Air supply is not needed for FiO ₂ = 100 Vol.%. <hr/>	Ensure pressure is less than 6 bar.
!!! Airway pressure high	The upper alarm limit for the airway pressure has been exceeded. The patient is "fighting" the ventilator, cough.	Check patient condition, check ventilation pattern, correct alarm limit if necessary.
	Ventilation hose buckled.	Check hose system and tube.
!!! Airway pressure low	Leaking cuff.	Inflate cuff and perform leak test.
	Leak or disconnection.	Check hose system for tight connections. Check that the expiration valve is properly engaged.
!!! Apnoea	Patient's spontaneous breathing has stopped.	Apply controlled ventilation.
	Stenosis	Check condition of patient. Check tube.
	Flow sensor not calibrated or faulty.	Calibrate flow sensor. Replace if necessary.
! Apnoea alarm off	Flow monitoring deactivated.	Reactivate flow monitoring, page 69, or ensure adequate external monitoring immediately.
	In NIV mode: Apnoea monitoring has been deactivated.	Re-set the upper alarm limit for apnoea monitoring to the required value. Note Instructions for Use of NIV mode.

Message	Cause	Remedy
!! Apnoea ventilation	Due to detected apnoea, the system has switched over automatically to mandatory ventilation.	Check ventilation procedure. Return to the original ventilation procedure with »Alarm Reset«. Check condition of patient. Check tube.
! ASB > 1.5 s	Only appears in paediatric mode. The ASB cycle has been switched off 3 times due to time limitation.	Test ventilation system for leaks.
!!! ASB > 4 s	Only appears in adult mode. The ASB cycle has been switched off 3 times due to time limitation.	Test ventilation system for leaks.
!! Check settings	Power interruption while setting a ventilation pattern or the alarm limits.	Check pattern of ventilation and alarm limits. Confirm message with key »Alarm Reset«.
!!! Device failure	Device faulty.	Call DrägerService.
! Evita Remote ?	The Remote Pad has not been identified correctly.	Remove Remote Pad. Confirm message with key »Alarm Reset«. Call DrägerService at the next opportunity.
! Evita Remote inop.	Key pressed on Remote Pad during self-test.	Confirm message with key »Alarm Reset«. Remove Remote Pad and reconnect. Ensure that no key is pressed on the Remote Pad.
	Remote Pad faulty.	Confirm message with key »Alarm Reset«. Remove Remote Pad. Call DrägerService at the next opportunity.
!! Execute device check	Device check not performed.	Perform device check, page 31. Confirm message with »Alarm Reset« key.
! Exp. hold interrupted	The »Exp. hold« key has been pressed for more than 15 seconds.	Release the »Exp. hold« key.
!!! Exp. valve inop.	Expiration valve not properly connected to socket.	Push expiration valve firmly into socket until it clicks into place.
	Flow sensor not calibrated or defective.	Calibrate flow sensor, page 67, replace if necessary.
	Expiration valve faulty.	Replace expiration valve.
! External Flow	Evita 2 dura calculates the externally supplied flow when monitoring correct functioning of the flow measurement.	Deactivate calculation of the external flow, see page 68.
!!! Fan failure	Fan failure.	Call DrägerService.
!!! FiO2 high	O2 sensor not calibrated.	Calibrate O2 sensor, page 66.
	Faulty mixer function.	Call DrägerService.
!!! FiO2 low	O2 sensor not calibrated.	Calibrate O2 sensor, page 66.
	Faulty mixer function.	Call DrägerService.
!!! Flow measurement inop.	Water in flow sensor.	Dry flow sensor.
	Flow sensor faulty.	Calibrate flow sensor, page 67, replace if necessary.
	Flow measurement malfunction.	Call DrägerService.

Message	Cause	Remedy
! Flow monitoring off	Flow monitoring is switched off.	Switch on flow monitoring again, as described on page 69, or immediately ensure an adequate external monitor function.
!!! Flow sensor ?	Flow sensor not fully inserted in rubber lip of expiration valve.	Insert flow sensor correctly.
!! Hard key xx failed	Key xx (e.g. »  «) can no longer be pressed.	Call DrägerService.
!!! High frequency	Patient is breathing at a high spontaneous frequency	Check condition of patient, check pattern of ventilation, correct alarm limit if necessary.
! Insp. hold interrupted	The » Insp. hold « key was held down longer than 15 seconds.	Release » Insp. hold « key.
!!! Insp/Exp cycle failure	The device does not deliver any gas.	Check the P _{max} /PEEP setting. Set an IPPV frequency of at least 4/min. Increase T _{Apnoea}  alarm time.
	Device faulty.	Call DrägerService.
!! Key xx overused ?	Key has been pressed several times in a short period (e.g. »  «).	Confirm message with key » Alarm Reset «. Contact DrägerService if this message reoccurs frequently.
!! Key overused ?	Due to very frequent key use, the screen contents of the display are repeatedly redrawn.	Confirm message with key » Alarm Reset «.
	Brief communication failure between the display processor and main processor.	Confirm message with key » Alarm Reset «. Contact DrägerService if this message reoccurs frequently.
! Leakage	The measured leakage minute volume MV _{leak} is 20 % higher than the minute volume measured on the expiration side.	Check that the hose connection is leakproof. Check that the tube is correctly fitted.
!!! Loss of data	Lithium battery discharged.	Call DrägerService.
! Malfunction fan	Temperature in machine too high.	Check fan function, clean cooling-air filter or call DrägerService.
! MEDIBUS COM. inop.	The connector of the MEDIBUS cable was unplugged during operation.	Plug the connector in again and secure it against disconnection with the two screws.
	MEDIBUS cable defective.	Use a new MEDIBUS cable.
	Interface defective.	Call DrägerService.
!!! Mixer inop.	Mixer malfunction. FiO ₂ can deviate considerably.	Immediately ventilate with separate manual ventilation device! Call DrägerService.
! Multi functional board inop.	The multi-functional board for operating the nurse call or Remote Pad is faulty.	Confirm message with key » Alarm Reset «. Call DrägerService at the next opportunity. The original ventilation functions of Evita 2 dura are not affected. Correct functioning of the nurse call or Remote Pad is not guaranteed, however: remove the nurse call and/or Remote Pad.

Message	Cause	Remedy
!! Multi functional board inop.	The multi-functional board for operating the nurse call or Remote Pad is faulty.	Confirm message with key »Alarm Reset«. Call DrägerService at the next opportunity. The original ventilation functions of Evita 2 dura are not affected. Correct functioning of the nurse call or Remote Pad is not guaranteed, however: remove the nurse call and/or Remote Pad.
!!! MV high	The minute volume has exceeded the upper alarm limit.	Check condition of patient, check pattern of ventilation, correct alarm limit if necessary.
	Flow sensor not calibrated or faulty.	Calibrate flow sensor, page 67, replace if necessary.
	Water in flow sensor.	Drain water trap in hose system. Dry flow sensor.
	Machine malfunction.	Call DrägerService.
!!! MV low	The minute volume has fallen below the lower alarm limit.	Check condition of patient, check pattern of ventilation, correct alarm limit if necessary.
	Stenosis.	Check condition of patient. Check tube.
	Leak in breathing system.	Establish leakproof breathing system.
	Flow sensor not calibrated or faulty.	Calibrate flow sensor, page 67, replace if necessary.
	Machine malfunction.	Call DrägerService.
!! Nebulisation interrupted	Only in paediatric mode. Nebulisation is only possible in pressure-controlled ventilation or with AutoFlow®.	Select the patient mode. Restart nebulisation. Acknowledge the alarm with »Alarm Reset«.
	Only in paediatric mode, only for ventilation with AutoFlow. Flow sensor not ready for measurement.	Switch on flow monitoring or calibrate flow sensor, page 51, or replace flow sensor or change mode. Restart nebulisation. Acknowledge the alarm with »Alarm Reset«.
! Nebulizer on	The medicament nebuliser is switched on, page 60.	Switch off the medicament nebuliser if necessary, page 60.
!!! O2 measurement inop.	O2 sensor provides invalid measured values.	Calibrate O2 sensor, page 66, replace if necessary.
	O2 measurement malfunction.	Call DrägerService.
! O2 monitoring off	O2 monitoring switched off.	Switch on O2 monitoring again, as described on page 66, or immediately ensure an adequate monitor function.
!!! O2 supply down	O2 supply pressure too low.	Make sure pressure is greater than 3 bar.
! O2 supply down	O2 supply pressure too low. O2 supply pressure is not required when FiO2 = 21 Vol.%.	Make sure pressure is greater than 3 bar.
!! O2 supply pressure high	O2 supply pressure too high.	Make sure pressure is less than 6 bar.

Message	Cause	Remedy
! O₂ supply pressure high	O ₂ supply pressure too high. O ₂ supply pressure is not required when FiO ₂ = 21 Vol.%. FiO ₂ = 21 Vol.%.	Make sure pressure is less than 6 bar.
!!! PEEP high	Expiratory system obstructed.	Check hose system and expiration valve.
	Expiratory resistance is increasing.	Check bacterial filter. Replace if necessary.
	Machine faulty.	Call DrägerService.
!!! PEEP valve inop.	Internal PEEP valve faulty.	Call DrägerService.
! Pressure limited	P _{max} pressure limit is active.	Check condition of patient, check pattern of ventilation, correct setting if necessary.
!!! Pressure meas. inop.	Fluid in expiration valve.	Replace expiration valve, page 92, then clean and dry.
	Pressure measurement malfunction.	Call DrägerService.
!!! Standby activated	Evita 2 dura has been switched to standby.	Confirm standby with »Alarm Reset« key.
!!! Temperature high	Breathing gas temperature higher than 40 °C.	Switch off humidifier.
!!! Temperature meas. inop.	Temperature sensor faulty.	Fit new temperature sensor, see page 22.
!!! Temperature sensor ?	Temperature sensor probe has been disconnected during operation.	Reconnect probe.
	Sensor cable broken.	Fit new temperature sensor.
!!! Tidal volume high	The upper alarm limit of the applied inspiratory tidal volume VT has been exceeded during three consecutive ventilation strokes.	Check condition of patient, check pattern of ventilation, correct alarm limit if necessary.
	Leak or disconnection.	Check that hose system connections are leakproof.
! Tidal volume high	The inspiratory tidal volume VT has exceeded the upper alarm limit.	Check condition of patient, check pattern of ventilation, correct alarm limit if necessary.
	Leak or disconnection.	Check that hose system connections are leakproof.
!!! Tube blocked	Evita 2 dura only applies a very small volume with each mechanical stroke, e.g. because the tube is blocked.	Check condition of patient, check tube.
	Patient "fights" against the mechanical strokes in pressure-controlled ventilation, so that the set inspiratory pressure volume is achieved with only a very small volume.	Check condition of patient, check machine settings.
!! Volume not constant	Due to pressure limit or time limit, the set tidal volume VT has not been applied.	Prolong inspiratory time »T _{insp} « Increase inspiratory flow »Flow« Increase pressure limit »P _{max} «. Press the »Alarm Reset« key to suppress the visual and acoustic alarm until the cause of the alarm is remedied.
!!! CO₂ measurement inop	CO ₂ sensor faulty.	Replace defective CO ₂ sensor.
	CO ₂ electronics in unit faulty.	Call Dräger Service.

Care

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Care

- Note the hospital hygiene regulations!

The ventilator must be cleaned and prepared after every patient.

To avoid all risk of infection by hospital staff and other patients, the ventilator must be disinfected and cleaned before it is used on the next patient (Use protective clothing, eye protection, etc.).

Removing parts

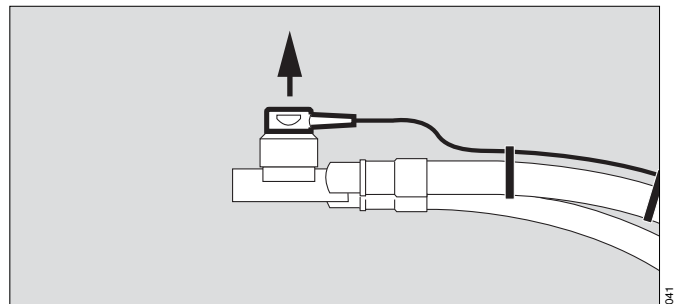
- Switch off the ventilator and humidifier, and remove their power plugs.
- Drain the water traps and ventilation hoses.
- Drain the water container of the humidifier.

Humidifier

- Dismantle in accordance with the specific Instructions for Use and prepare for disinfecting/sterilising.

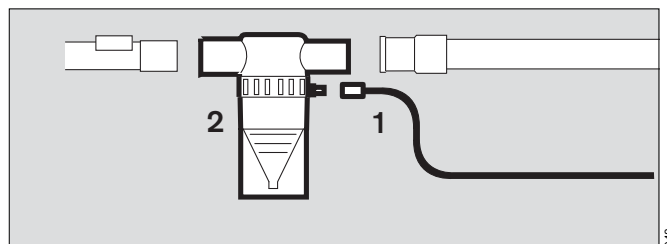
Temperature sensor (option)

- Remove the temperature sensor from the Y-piece – do not tug on cable. Remove the sensor probe from the back panel of Evita 2 dura.
- The temperature sensor is designed for wipe-disinfection. **The temperature sensor must not be placed in a cleaning and disinfection machine or disinfectant bath.**

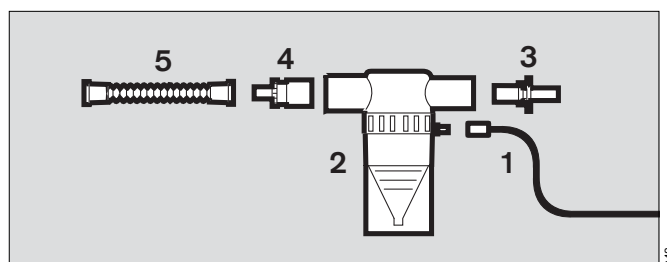


Medicament nebuliser (option)

- 1 Disconnect nebuliser hose from medicament nebulizer and from the port on the unit.
- 2 Disconnect the medicament nebuliser from the adult hose system
or



- 2 from the paediatric hose system.
 - 3 Remove catheter connector (ISO cone Ø 15/Ø 11) from the inlet.
 - 4 Remove the adapter (ISO cone Ø 22/Ø 11) from the outlet.
 - 5 Remove the corrugated hose from the adapter.
- Dismantle the medicament nebuliser in accordance with its Instructions for Use.
 - Prepare the individual parts of the medicament nebuliser and the adapter parts for cleaning and disinfection in a cleaning and disinfection machine.



Ventilation hoses

- Remove ventilation hoses from the device ports.
- Remove the water traps from the ventilation hoses. Remove the water containers from the water traps.
- Prepare the ventilation hoses, water traps and their collecting jars and the Y-piece for cleaning and disinfection in a cleaning and disinfection machine.

Flow-Sensor

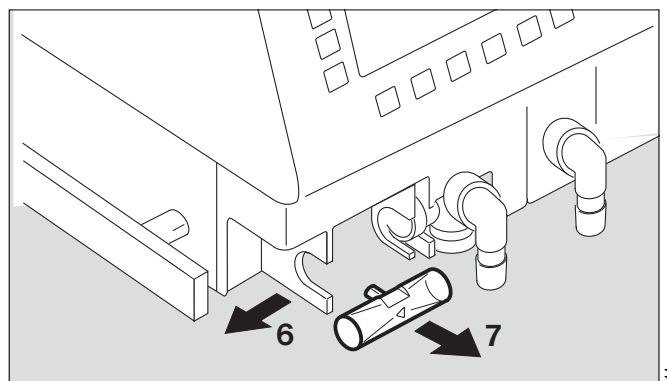
- 6 Push the flow sensor to the left as far as it will go and
- 7 pull out.

The flow sensor cannot be disinfected/cleaned in a machine, nor can it be sterilised by the hot steam method.
See page 95 for cleaning instructions of the flow sensor.

- Disinfect flow sensor for about 1 hour in 70 % ethanol solution.

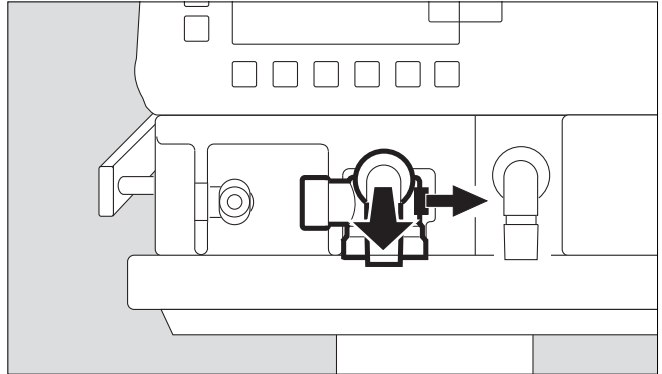
Expose the sensor to air for at least 30 minutes to allow the alcohol to evaporate. Otherwise the sensor could be damaged beyond control by ignition of any residual alcohol during calibration.

- The flow sensor may be re-used as long as calibration can be carried out successfully.



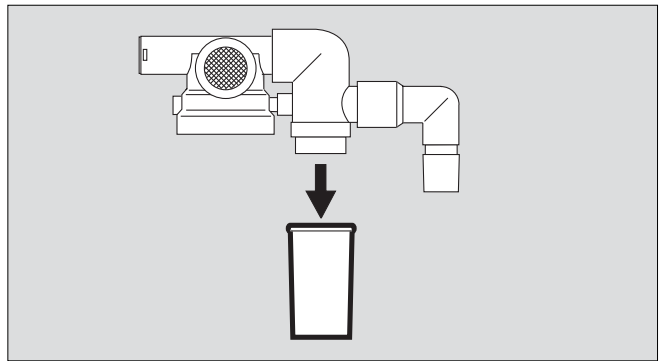
Expiration valve

- Push the catch to the right, pulling off the expiration valve at the same time.



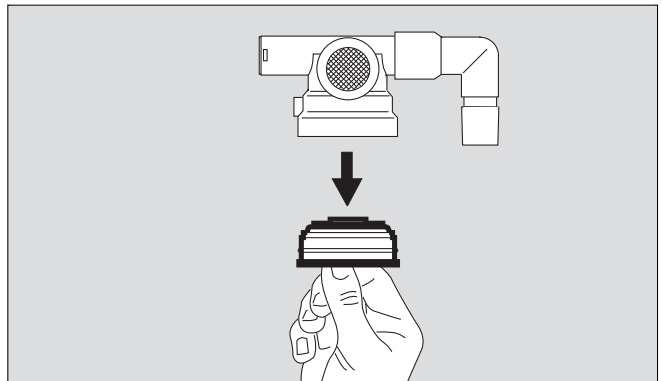
If the expiration valve has an optional water trap:

- Remove the collecting jar.



Only strip down the expiration valve if badly soiled:

- Unscrew stopper by hand and remove together with the diaphragm.
- Do not disassemble the expiration valve any further.
- Prepare the expiration valve for cleaning and disinfection in a cleaning and disinfection machine, then
- for sterilisation by the hot steam method.



Disinfecting/Cleaning

Use surface disinfectants. For surface compatibility, use disinfectants based on:

- aldehydes,
- quaternary ammonium compounds.

To avoid the possibility of damage to material, do not use any disinfectants based on:

- alkylamine-based compounds
- phenol-based compounds,
- halogen-releasing compounds,
- strong organic acids,
- oxygen-releasing compounds.

For users in the Federal Republic of Germany, we recommend that only disinfectants on the current DGHM list are used (DGHM: German Society for Hygiene and Microbiology). The DGHM list (published by mhp-Verlag, Wiesbaden) also classifies each disinfectant by its active agents.

For countries where the DGHM list is not available, we recommend the types of disinfectant given above.

Disinfectants often contain – besides their main active agents – additives that can also damage materials.

If in doubt, ask the supplier/manufacturer of the disinfectant/cleaning agent.

Parts must not be sterilised in ethylene oxide – health hazard!

To avoid all risk of infection by hospital staff and other patients, the ventilator must be disinfected and cleaned before it is used on the next patient (Use protective clothing, eye protection, etc.).

Ventilator without ventilation hoses, gas supply hoses, temperature sensor

Wipe disinfect

- e.g. with Buraton 10 F or Terralin (Schülke & Mayr, Norderstedt, Germany). Comply with the manufacturer's instructions.

Cooling air filter, room air filter

- Filters must be cleaned or replaced when soiled or at the latest after 4 weeks, see page 98.

Ventilation hoses, water traps and their collectors, Y-piece, expiratory valve (or in the event of severe soiling, its individual components)

- Disinfect in a moisture saturated environment at 93 °C (200 °F) for 10 minutes using a cleaning and disinfecting machine. **Use detergent only.**
- After disinfecting with moist heat, we recommend that the expiratory valve or its disassembled components be autoclaved **at 134 °C (273 °F)** to remove any remaining liquid in the pressure measuring canal in the block.

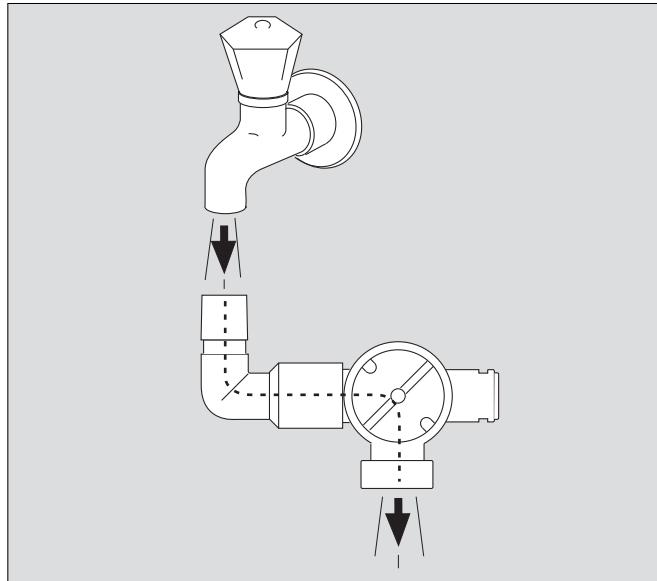
Alternatively:

If a cleaning and disinfection machine is not available:

- Bath disinfect, e.g. with Sekusept from Henkel. Comply with the manufacturer's instructions.
- Then rinse with clean water, preferably from a soft water supply. Shake water out thoroughly, and leave the products to dry.

Expiration valve and its individual parts after disinfection

- Rinse thoroughly with clear water, preferably from a soft water supply. Shake water out thoroughly.
- After rinsing thoroughly, dry expiration valve.
- After drying, sterilise in hot steam at 134 °C, otherwise liquid may remain in the pressure measuring line and impair correct functioning.



Ventilation hoses, water traps and their collectors, Y-piece, expiratory valve, temperature sensor

- Can be autoclaved at 134 °C (273 °F).

Humidifier

- Prepare in accordance with separate Instructions for Use.

Care list for Evita 2 dura Intensive Care Ventilator

Applicable for use with non-infectious patients.

For infectious patients, all parts that conduct breathing gas must be additionally sterilised after disinfecting and cleaning.

The parts in contact with breathing gas and listed below can be sterilised in hot steam at 134 °C. Refer to the column "Sterilising".

The below/above table contains the manufacturers recommended guidelines. Please note that these are merely guidelines and are not intended to take precedence over the hospitals hygiene/infection control protocols!!

Part Reusable components	How often Recommended cleaning intervals	How			Sterilising Steam 134 °C 10 minutes
		Disinfecting and cleaning			
		Cleaning and disinfection machine, 93 °C 10 minutes	Wiping	Bath immersion	
Evita 2 dura basic device	per patient	no	outside	no	no
Trolley Hinged arm Medical gas hoses	per patient	no	outside	no	no
Ventilation hoses, Y-piece, Water traps, Collecting jars	per patient/ weekly	yes	no	possible	yes
Expiration valve	per patient/ weekly*	yes**	no	possible**	yes
Temperature sensor	per patient/ weekly*	no	yes	no	yes
Flow sensor	daily	no***	outside	possible***	no

* Nebulisation may lead to formation of more extensive deposits necessitating more frequent replacement.

** After disinfecting/cleaning: sterilise at 134 °C. Otherwise risk of malfunction due to residual liquid in pressure measuring line.

*** Special treatment, see page 91.

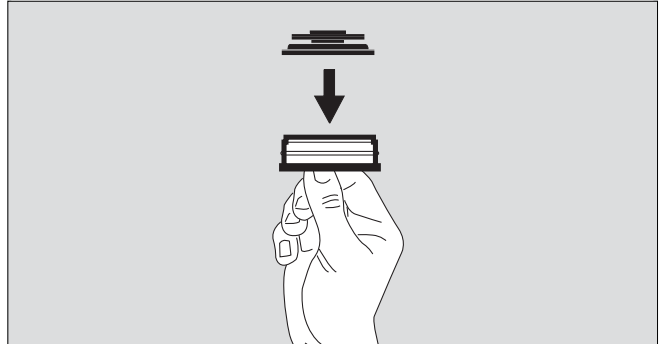
Assembling

Mounting the expiration valve

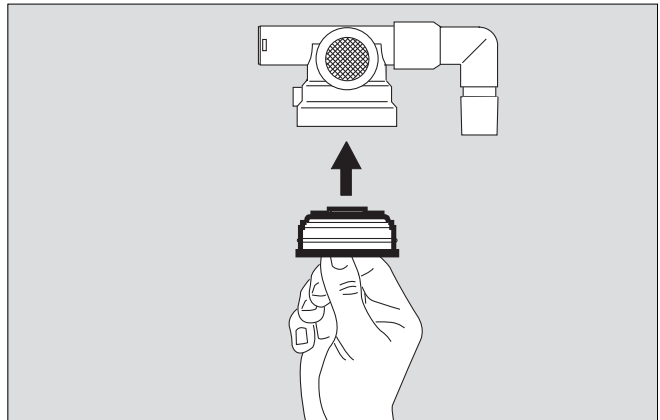
The parts must be entirely dry to prevent malfunctioning.

Be careful to fit the diaphragm in the correct position.

- Hold stopper by the flange and place diaphragm on the collar of the stopper.

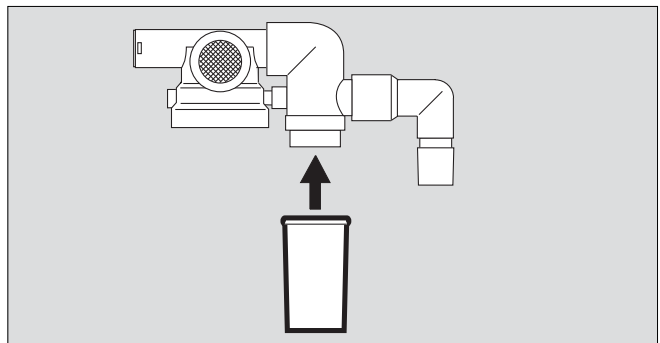


- Insert stopper with diaphragm on top into the housing from below and screw in tightly.



If the expiration valve has an additional water trap:

- Fit collecting jar.



Medication nebuliser

- Assemble in accordance with separate Instructions for Use.
- Installation, see page 61.

Breathing gas humidifier

- Assemble in accordance with separate Instructions for Use.

Before re-using on patient

- Assemble machine as described under "Preparing for use" on page 18 onwards.
- Carry out checks to ensure readiness for operation, see "Device Check" on page 30 onwards.

Maintenance Intervals

Clean and disinfect equipment and/or components before any maintenance procedures and/or before returning for repair!

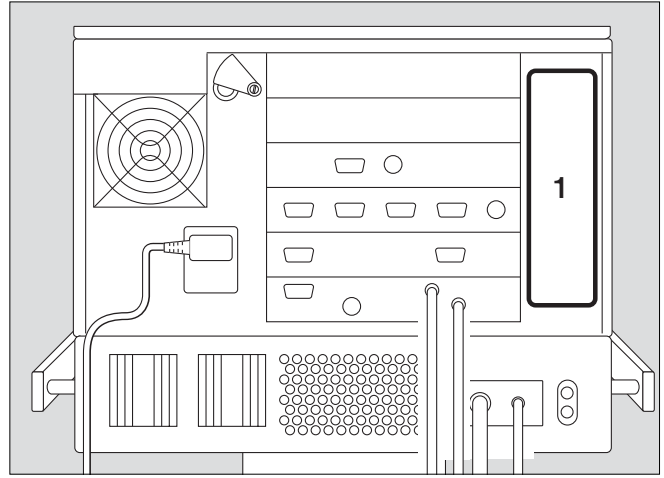
Note the maintenance intervals specified in the respective Instructions for Use of the options installed!

O ₂ sensor capsule	Replace when the following message is displayed: »O ₂ measurement inop« and when calibration is no longer possible. Spent O ₂ sensors can be returned to Dräger for disposal.
Ambient-air filter	Clean or replace after 4 weeks, see page 98.
Cooling-air filter	Replace after 1 year. Dispose of with normal domestic waste.
Filters in the compressed gas inlets	To be replaced by trained service personnel every 2 years.
Lithium battery for data protection	To be replaced by trained service personnel every 2 years. Disposal: see page 99.
Real-time clock	To be replaced by trained service personnel every 6 years. Disposal: page 99.
Pressure reducer	To be replaced every 6 years by DrägerService.
Equipment inspection and service	Every 6 months by trained service personnel.

Clean or replace cooling air filter

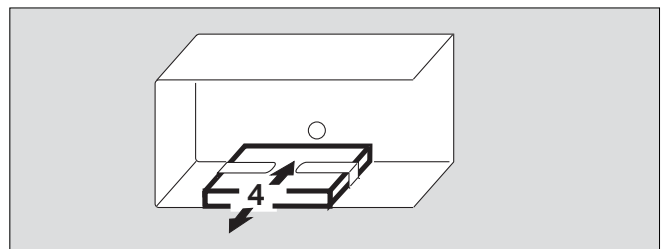
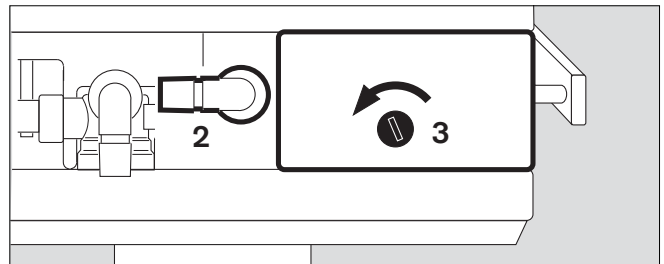
- Filter must be cleaned or replaced when soiled or at the latest after 4 weeks.
Replace after 1 year at the latest.
- 1 Remove cooling-air filter from its slot on the back of machine.
- Replace or clean in warm water with added detergent; dry well.
- Insert cooling-air filter in slot, taking care not to crease it.
- Dispose of used cooling-air filter with domestic waste.

Note the Instructions for Use of the option when using the (optional) DC power supply MB.



Removing and reinserting ambient-air filter

- Filter must be cleaned or replaced when soiled or at the latest after 4 weeks.
Replace filter every year.
 - 2 Swivel port to the left.
 - 3 Unscrew the screw (e.g. with a coin), and remove the protective cover.
-
- 4 Remove the ambient-air filter from the protective cover.
 - Push the new/cleaned ambient-air filter under the lugs.
 - Replace protective cover, and tighten screw with a coin.
 - Dispose of used ambient-air filter with domestic waste.



Correct disposal of batteries and O₂ sensors

Batteries and O₂ sensors:

- Do not incinerate or throw in fire; risk of explosion!
- Do not open using force; risk of corrosion!
- Do not re-charge batteries.

Batteries must be disposed of as special waste:

- Information may be obtained from the local environmental and public health authorities or from approved waste disposal companies.

O₂ sensors can be returned to Dräger.

Correct disposal of apparatus

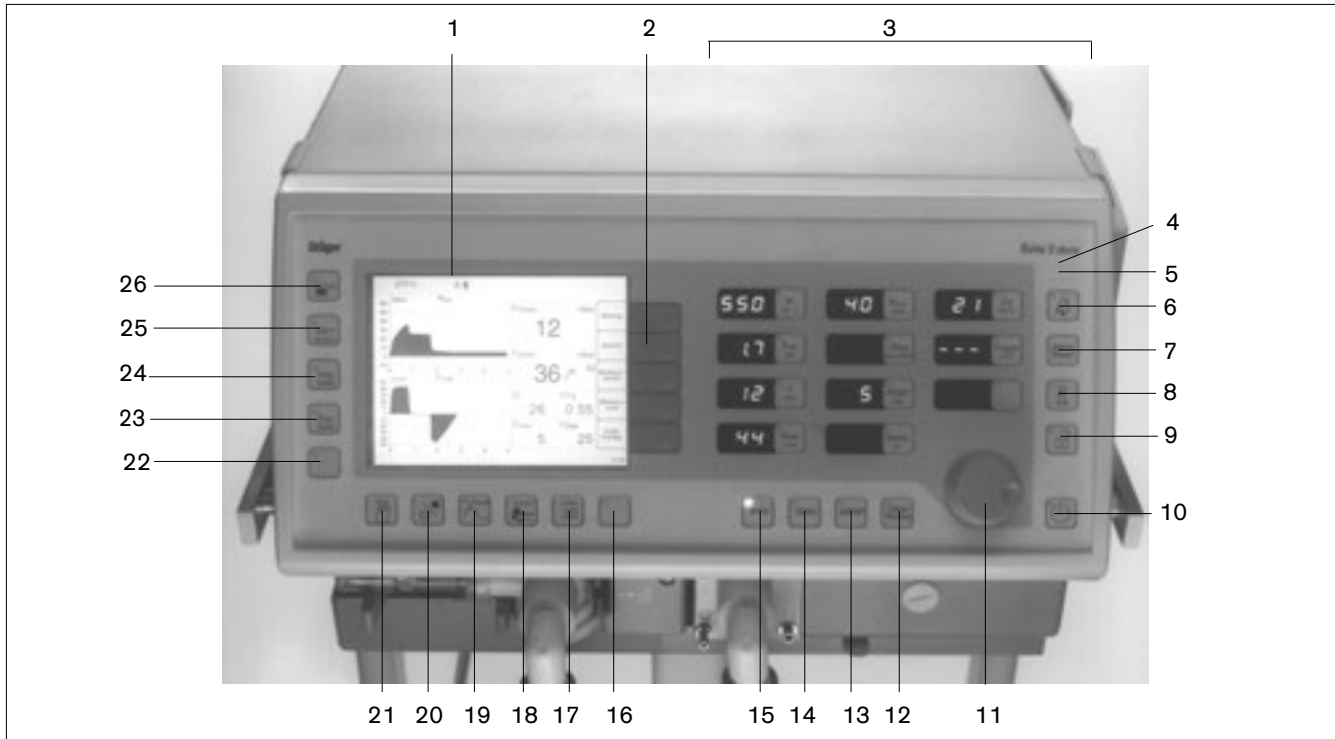
- at the end of its useful life
- After contacting the competent waste disposal company, hand over Evita 2 dura for appropriate disposal. The applicable legal regulations must be observed.

What's what

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Control unit	102
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Back panel, with DC power supply MB	105

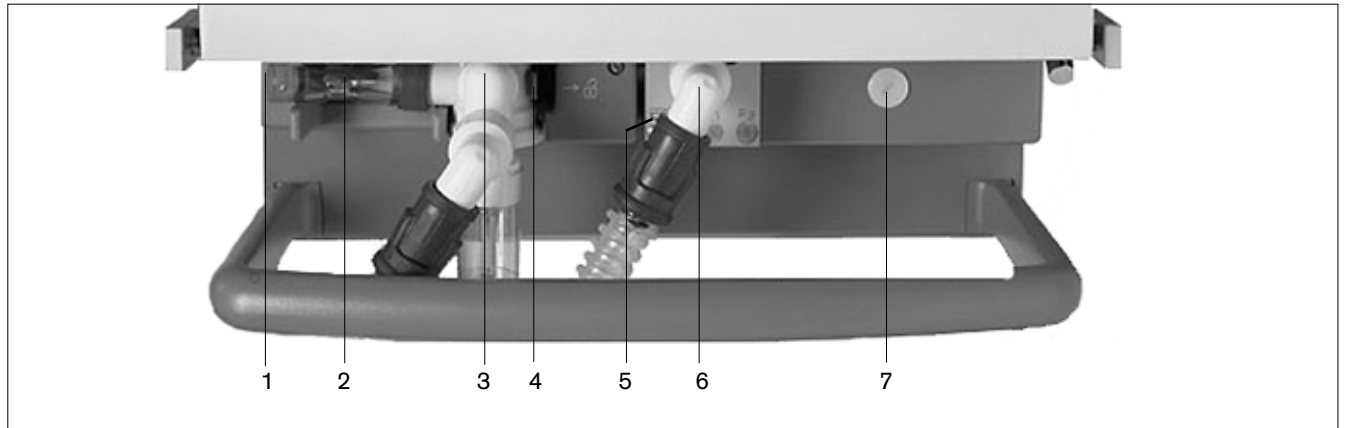
What's what

Control unit



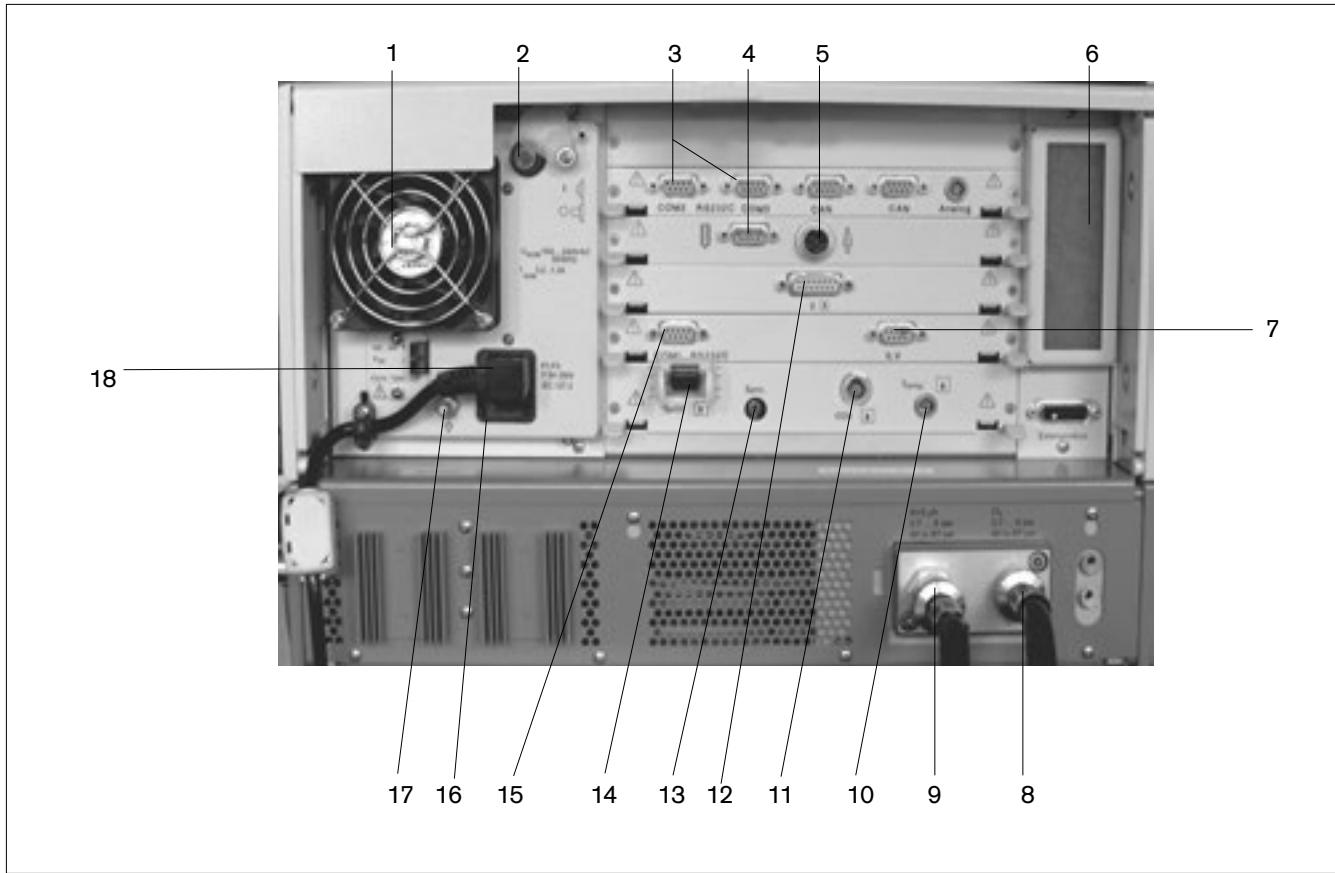
- | | |
|---|---|
| <ul style="list-style-type: none"> 1 Screen for displaying application-specific screen pages. 2 Menu keys for displaying the application-specific screen pages. 3 Parameter keys with displays indicating their settings – for setting the ventilation parameters. 4 Red LED to indicate warnings 5 Yellow LED to indicate cautions and advisory messages 6 »« key for suppressing the audible alarm tone for 2 minutes. 7 »Alarm Reset« key for acknowledging alarm messages 8 »« key for calling up information and help on settings 9 »« key ("lock") for protecting the ventilation parameters and ventilation mode against unauthorised modification. 10 »« key ("standby") for changing between ventilation and standby 11 Central "turn and press" rotary knob for selecting and confirming settings 12 »Other Modes« key for using other ventilation modes programmed on-screen | <ul style="list-style-type: none"> 13 Key for the BIPAP ventilation mode 14 Key for the SIMV ventilation mode 15 Key for the IPPV ventilation mode 16 Key for future functions 17 »Values « key for changing the displayed value set 18 »Curves « key for changing the displayed curve pair 19 »Freeze « key for freezing curves 20 »/« key for switching the screen brightness between bright/dark 21 »Print « key for manual printer logging 22 Key for future functions 23 »Exp. hold« key for manually extending the expiration phase 24 »Insp. hold« key for manual inspiration 25 »O₂ ↑ Suction« key for bronchial suction 26 »« key for switching the pneumatic medicament nebuliser on/off |
|---|---|






Front connections



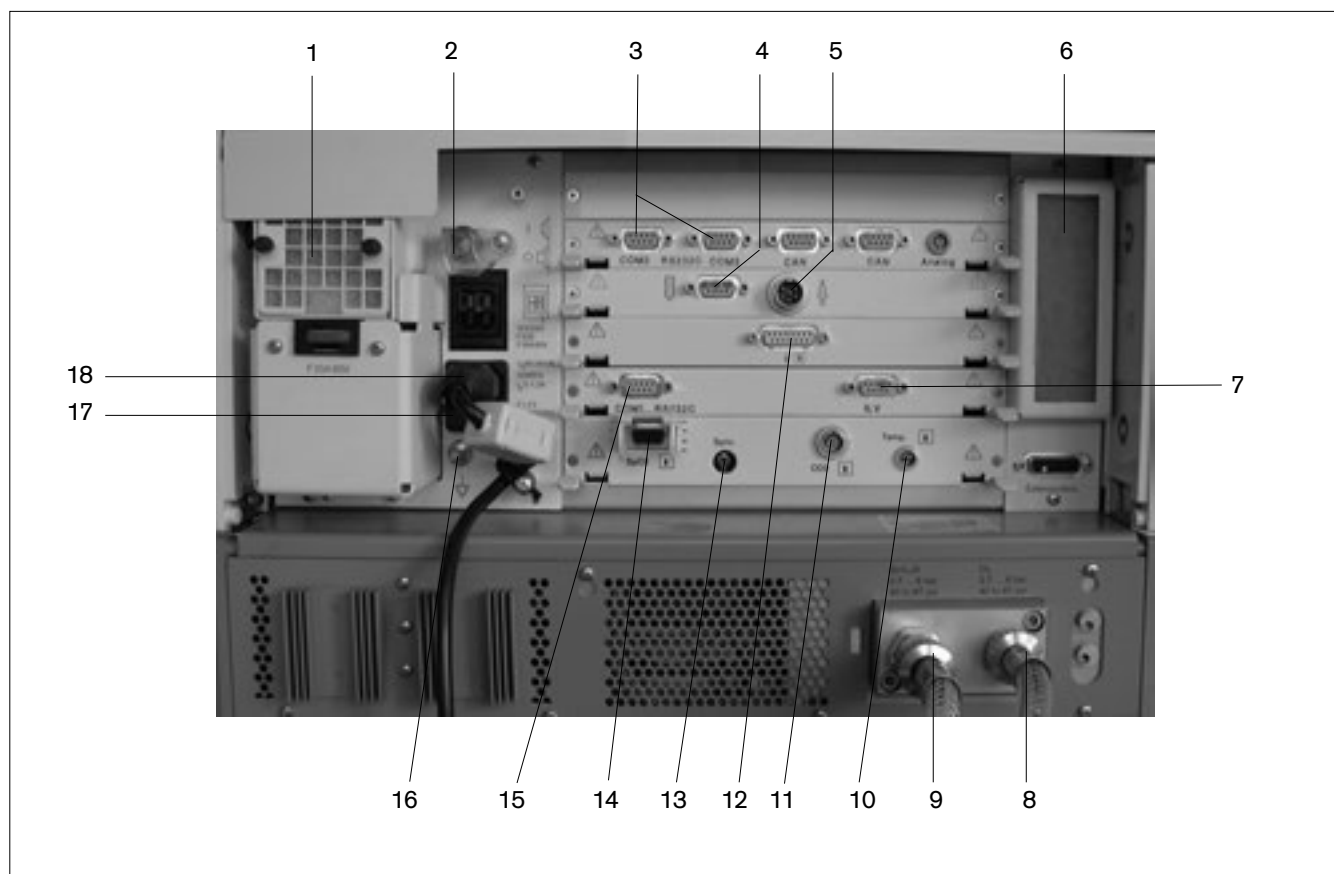
- 1 Gas outlet (EXHAUST – NOT FOR SPIROMETERS)
- 2 Flow sensor
- 3 Expiration valve with expiration port (GAS RETURN)
- 4 Latch for expiration valve
- 5 Nebuliser port
- 6 Inspiratory port (GAS OUTPUT)
- 7 Locking screw for protective cover
(behind it: O₂ sensor and ambient-air filter)






Back panel



- | | |
|---|--|
| <p>1 Fan</p> <p>2 Power switch with protective flap</p> <p>3 »COM2«, »COM3« sockets for RS232 and analogue interfaces (optional)</p> <p>4 Connection »« for Remote Pad, optional</p> <p>5 Connection »« for nurse call, optional</p> <p>6 Cooling-air filter</p> <p>7 ILV socket, optional</p> <p>8 Connection for oxygen</p> <p>9 Connection for medical air</p> <p>10 »Temp. « socket for temperature sensor</p> | <p>11 »CO2 « socket for CO2 sensor, optional</p> <p>12 Connection for neonate flow sensor (optional)</p> <p>13 »Sync.« socket for C-Lock-ECG synchronisation for optional SpO2 measurement, optional</p> <p>14 »SpO2 « socket for functional SpO2 measurement, optional</p> <p>15 »COM1 RS232C« socket for RS232 interface, e.g. for printer</p> <p>16 Mains fuses</p> <p>17 Earth connection</p> <p>18 Connector for power cord</p> |
|---|--|

Back panel, with DC power supply MB



- | | |
|--|---|
| <p>1 Fan</p> <p>2 Power switch with protective flap</p> <p>3 »COM2«, »COM3« sockets for RS 232 and analogue interfaces (optional)</p> <p>4 Connection »« for Remote Pad, optional</p> <p>5 Connection »« for nurse call, optional</p> <p>6 Cooling-air filter</p> <p>7 ILV socket</p> <p>8 Connection for oxygen</p> <p>9 Connection for medical air</p> <p>10 »Temp. « socket for temperature sensor</p> | <p>11 »CO2 « socket for CO2 sensor, optional</p> <p>12 Connection for neonate flow sensor (optional)</p> <p>13 »Sync.« socket for C-Lock-ECG synchronisation for optional SpO2 measurement, optional</p> <p>14 »SpO2 « socket for functional SpO2 measurement, optional</p> <p>15 »COM1 RS232C« socket for RS 232 interface, e.g. for printer</p> <p>16 Earth connection</p> <p>17 Mains fuses</p> <p>18 Connector for power cord</p> |
|--|---|

7452002

Technical Data

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Technical Data

Environmental conditions

In operation

Temperature	10 to 40 °C
Atmospheric pressure	700 to 1060 hPa
Rel. humidity	0 to 90 %, without condensation

In storage

Temperature	-20 to 60 °C
Atmospheric pressure	500 to 1060 hPa
Rel. humidity	0 to 100 %, without condensation

Settings

Ventilation modes

IPPV/IPPV_{Assist},
SIMV, SIMV/ASB
MMV, MMV/ASB
BIPAP/ASB
BIPAP_{Assist}
CPAP/ASB

Ventilation frequency f

0 to 100/min

Inspiration time T_{insp}

0.1 to 10 s

Inspiration time V_T

Paediatrics	0.02 to 0.3 L, BTPS*
Accuracy	±10 % of set value, or ±10 mL, whichever is greater.
Adults	0.1 to 2.0 L, BTPS*
Accuracy	±10 % of set value, or ±25 mL, whichever is greater.

Inspiratory Flow

Paediatrics	6 to 30 L/min
Adults	6 to 120 L/min

Inspiratory pressure P_{insp}

0 to 80 mbar

Inspiratory pressure limit P_{max}

0 to 100 mbar

O₂ concentration

21 to 100 Vol.%

Accuracy	±5 % of set value, or ±2 Vol.%, whichever is greater.
----------	---

Positive end-expiratory pressure PEEP or intern. PEEP

0 to 35 mbar

Trigger sensitivity

0.3 to 15 L/min

Pressure assist PASB

0 to 80 mbar

Rise time for pressure support (Ramp)

0 to 2 s

I:E

1:9.5 to 4:1 (can only be configured in default settings)

* BTPS = Body Temperature, Pressure, Saturated.
Measured values with reference to the conditions of the patient lung: Body temperature 37 °C, steam-saturated gas, ambient pressure.

Performance data

Control principle	Time-cycled, volume-constant, pressure-controlled
Intermittent PEEP frequency	2 cycles every 3 minutes
Medicament nebulisation	for 30 min
Bronchial suction	
Disconnection detection	automatic
Reconnection detection	automatic
Oxygen enrichment	max. 3 minutes
Active suction phase	max. 2 minutes
Final oxygen enrichment	2 minutes
Valve response time T _{0...90}	≤5 ms
Supply system for spontaneous breathing and ASB	adaptive CPAP system with high initial flow
Max. flow rate	2 L/s in 8 ms
Max. inspiratory flow	180 L/min
Equipment compliance (with Aquapor humidifier and patient tubing system for adults)	≤2 mL/mbar
Inspiration resistance	≤2.3 mbar/L/s
Expiration resistance	≤3.8 mbar/L/s
Equipment compliance (with Fisher & Paykel MR 730 humidifier and tubing system K for paediatric use)	≤1 mL/mbar
Inspiration resistance	≤4.1 mbar/L/s
Expiration resistance	≤4.1 mbar/L/s
Additional functions	
Inspiratory relief valve	Opens if medical air supply fails (pressure <1.2 bar), enables spontaneous breathing with filtered ambient air
Safety valve	Opens the breathing system at 100 mbar.

Measured value displays

Airway pressure measurement	
Max. airway pressure	P _{peak}
Plateau pressure	P _{plat}
Pos. end-exp. pressure	PEEP
Mean airway pressure	P _{mean}
Min. airway pressure	P _{min}
Range	0 to 99 mbar
Resolution	1 mbar
Accuracy	±2 mbar

Technical Data

Measured value displays

O₂ measurement in main flow (inspiratory side)

Inspiratory O₂ concentration FiO₂

Range	15 to 100 Vol.%
Resolution	1 Vol.%
Accuracy	±3 Vol.%

Flow-Messung

Minute Volume MV

Spontaneously breathed minute volume MV_{spn}

Range	0 to 99 L/min, BTPS*
Resolution	0.1 L/min
Accuracy	±8 % of measured value
To...90	approx. 35 s

Tidal volume VT_e

Spontaneously breathed tidal volume VT_{spn}

Range	0 to 3999 mL, BTPS*
Resolution	1 mL
Accuracy	±8 % of measured value

Tidal volume VT_{ASB}

Inspiratory tidal volume during an ASB stroke

Range	0 to 3999 mL, BTPS*
Resolution	1 mL
Accuracy	±8 % of measured value

Frequency Measurement

Breathing frequency f_{tot}

Spontaneous breathing frequency f_{spn}

Range	0 to 150 /min
Resolution	1 /min
Accuracy	±1 /min
To...90	approx. 35 s

Breathing gas temperature measurement

Range	18 to 51 °C
Resolution	1 °C
Accuracy	±1 °C

* Body Temperatur, Pressure, Saturated.
Measured values relating to the conditions of the patient lung: Body temperature 37 °C, steam-saturated gas, ambient pressure.

Computed value displays

Compliance C

Range	0.7 to 200 mL/mbar
Resolution	
Range 0.7 to 99.9 mL/mbar	0.1 mL/mbar
Range 100 to 200 mL/mbar	1 mL/mbar
Accuracy	±20 % of measured value *

Resistance R

Range	3 to 200 mbar/L/s
Resolution	
Range 3 to 99.9 mbar/L/s	0.1 mbar/L/s
Range 100 to 200 mbar/L/s	1 mbar/L/s
Accuracy	±20 % of measured value **

Leakage minute volume MV_{leak}

Range	0 to 99 L/min, BTPS
Resolution	0.1 L/min or for values less than 0.1 L/min: 0.01 L/min
Accuracy	±18 % of measured value
T _{0...90}	approx. 35 s

Curve displays

Airway pressure P _{aw} (t)	-10 to 100 mbar
Flow (t)	-150 to 180 L/min
Volume V (t)	0 to 2000 mL

Rapid-Shallow-Breathing RSB

Range	0 to 9999 1/(min x L)
Resolution	1/(min x L)
Accuracy	see measurement of V _T and f

Negative Inspiratory Force NIF

Range	-45 to 0 mbar
Resolution	1 mbar
Accuracy	±2 mbar

* C-values may be considerably falsified as spontaneous breathing increases; compliance with the measuring accuracy therefore cannot be guaranteed for spontaneous breathing.

** R-values may be considerably falsified as spontaneous breathing increases; compliance with the measuring accuracy therefore cannot be guaranteed for spontaneous breathing.

Monitoring

Expiratory minute volume **MV**

Upper alarm limit alarm

Setting range

when MV exceeds the upper alarm limit.

0.1 to 1 L/min in 0.01 L/min increments

1 to 41 L/min in 0.1 L/min increments

Lower alarm limit alarm

Setting range

when MV falls below the lower alarm limit.

0.01 to 1 L/min in 0.01 L/min increments

1 to 40 L/min in 0.1 L/min increments

Airway pressure **Paw**

Upper alarm limit alarm

Setting range

when the "Paw high" value is exceeded.

10 to 100 mbar

Lower alarm limit alarm

when the value "PEEP +5 mbar" (coupled with the PEEP set value) is not exceeded for at least 96 ms in 2 successive ventilation strokes.

Insp. O₂ concentration **FiO₂**

Upper alarm limit alarm

Lower alarm limit alarm

Range

if FiO₂ exceeds the upper alarm limit for at least 20 seconds.

if FiO₂ falls below the lower alarm limit for at least 20 seconds.

both alarm limits are automatically allocated to the set value:

less than 60 Vol.%, ±4 Vol.%

60 Vol.% and over: ±6 Vol.%

Insp. breathing gas temperature

Upper alarm limit alarm

when temperature reaches 40 °C.

(Evita 2 dura can also be used without temperature sensor if the sensor is not connected on switching on).

Tachypnoea monitoring **f_{spn}**

Alarm

when the spontaneous breathing frequency is exceeded during spontaneous breathing.

Adjustment range

5 to 120/min

Volume monitoring **VT_i**

Lower alarm limit alarm

if the set tidal volume VT (coupled with the set value VT) has not been supplied.

Upper alarm limit alarm

if the applied tidal volume exceeds the value of the alarm limit, inspiration is interrupted and the expiration valve is opened.

Adjustment range

21 to 4000 mL

Apnoea alarm time **T_{Apnoea}**

Alarm

if no breathing activity is detected.

Adjustment range

5 to 60 s, adjustable in 1 second steps.

Operating data

Mains power connection

100 V to 240 V 50/60 Hz



Current input

at 230 V

max. 1.3 A

at 100 V

max. 3.2 A

Power consumption	typically approx. 125 W
Machine fuses	
Range 100 V to 240 V	F 5 H 250 V IEC 127-2 (2x)
Protection class	
Machine	Class I
CO ₂ sensor (sensor connected)	Type BF 
Temperature sensor AWT 01 (sensor connected)	Type BF 
Gas supply	
O ₂ gauge pressure	3 bar – 10 % to 5.5 bar +10 % at 60 L/min (peak flow 200 L/min)
O ₂ connection thread	M 12 x 1, female
Air gauge pressure	3 bar – 10 % to 5.5 bar +10 % at 60 L/min (peak flow 200 L/min)
Air connection thread	M 20 x 1.5, male
Dew point	5 °C below ambient temperature
Oil concentration	<0.1 mg/m ³
Particle size	Dust-free air (filtered with filter size <1 µm)
Gas consumption of control system	Medical air or O ₂ approx. 3.5 L/min
Output for pneumatic medicament nebuliser	Medical air or O ₂ , max. 2.25 bar, max. 11 L/min
Automatic gas switch-over	if one gas fails (inlet pressure <1.5 bar) the device switches to the other gas.
Sound pressure (for free-field measurement over a reflecting surface)	max. 47 dB (A)
Dimensions (W x H x D)	
Basic machine	530 x 290 x 450 mm
Machine with trolley	580 x 1335 x 660 mm
Weight	
Basic machine	approx. 27 kg
Basic machine with trolley incl. cabinet 8H	approx. 69 kg

Machine outputs

Digital output	Output and reception via an RS 232 C interface
COM 1	LUST protocol Baudrate: 1200, 2400, 4800, 9600, 19200 baud Data bits: 7 Parity: even Stop bits: 1 MEDIBUS protocol Baudrate: 1200, 2400, 4800, 9600, 19200 baud Data bits: 8 Parity: even, odd, no Stop bits: 1 or 2 (19200 baud are required for transmission of high-speed data, e.g. for flow curve) Printer protocol HP Deskjet, series 500 Baudrate: 1200, 2400, 9600, 19200 baud Data bits: 8 Parity: no Stop bits: 1

Technical Data

Materials used

Cable length	Up to 15 m
Load impedance	3000 to 7000 Ω
Signal level (at load impedance 3000 to 7000 Ω)	
Low	Between 3 and 15 V
High	Between -3 and -15 V
Electrical isolation	Port COM 1 is electrically isolated from the machine electronics. The test voltage for the electrical isolation equals 1500 V.
Pin assignment	Pin 2 RxD Pin 3 TxD Pin 5 GND Connector housing Machine housing
Digital output	Output for independent lung ventilation (ILV)
Digital output (optional)	for output and reception via two RS 232 C interfaces
Digital output (optional)	for output and reception via a CAN interface
Analogue output (optional)	for output of analog data
Electromagnetic compatibility (EMC) (conforming to European Directive 89/336/EEC)	Tested in accordance with EN 60601-1-2
Classification as per EC Directive 93/42/EEC Annex IX	II b
UMDNS-Code Universal Medical Device Nomenclature System – Nomenclature for medical products	17-429

Materials used

Part	Appearance	Material
Ventilation hose	milky, transparent	silicone rubber
Water traps	yellow, transparent	polysulphone
Y-piece with connector for temperature measurement	milky, transparent	polysulphone
Expiration valve housing, closure	yellow, transparent	silicone rubber
Diaphragm	white	polyamide
CO ₂ cuvette	whitish and grey	silicone rubber and aluminium
Temperature sensor/cable	yellow, transparent	polysulphone with glass windows
CO ₂ sensor/cable	milky/green or blue	silicone rubber
	grey/grey	polyurethane

Description

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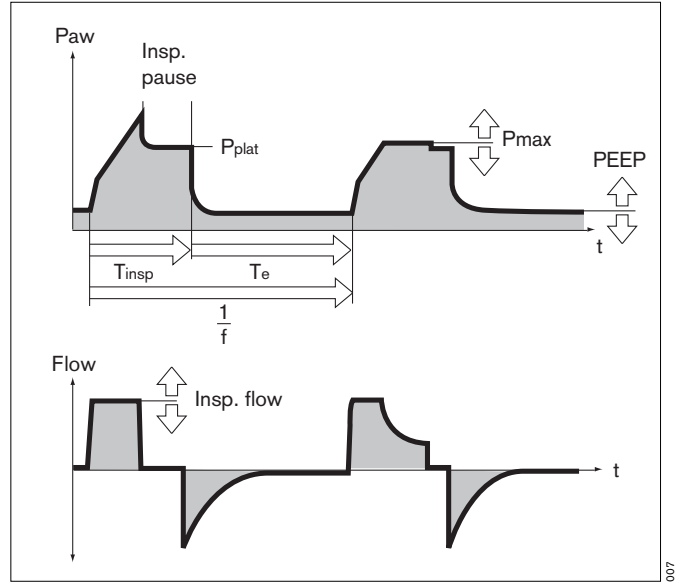
Description

Ventilation Modes

Volume-controlled ventilation with PLV

Classic volume-constant mandatory ventilation stroke

In mandatory ventilation strokes, the »Flow« parameter restricts the inspiration flow. If the inspiration flow is so high that the set tidal volume V_T is attained before the inspiration time T_{insp} has fully elapsed, the inspiration valve closes, and the supply of breathing gas stops. The expiration valve remains closed until the end of the inspiration time T_{insp} . This phase, the inspiratory pause, can be identified as the plateau P_{plat} in the curve $P_{aw}(t)$.



Manual pressure limiting with P_{max}

Evita 2 dura can prevent peaks of pressure, while maintaining the set tidal volume V_T , by means of the pressure limit P_{max} . The tidal volume V_T remains constant as long as a pressure plateau P_{plat} is still detectable and the flow curve shows a brief pause of zero flow between inspiration and expiration.

Evita 2 dura performs this function by reducing the inspiratory flow on reaching the set P_{max} value. If the tidal volume V_T can no longer be attained with the selected pressure P_{max} , due to reduced compliance, the alarm "Volume not constant" is automatically generated. Manual pressure limiting can be performed with all Evita models.

Sigh (intermittent PEEP)

"Sigh", in the form of intermittent PEEP, is operative in IPPV, IPPV_{Assist} and ILV modes.

The purpose of the expiratory sigh during ventilation is to open collapsed areas of the lung, or to keep open "slow" areas of the lung.

Since atelectatic alveoli have a longer time constant – also caused by obstructed bronchioles – increased airway pressure maintained over a longer period is required to open them.

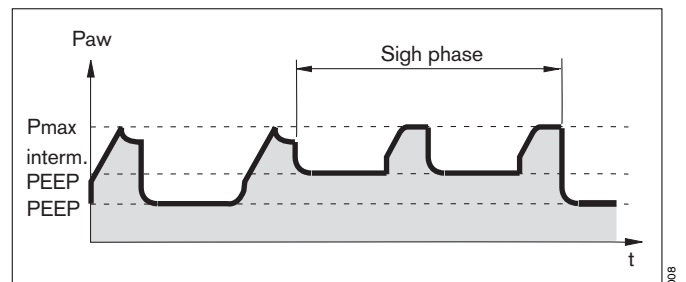
In many cases, the sigh function is achieved by increasing the ventilation stroke; however, due to the short time available, this form of sigh only marginally improves the filling of the "slow" alveoli.

In Evita 2 dura, the sigh operates during expiration with an intermittent PEEP for two ventilation strokes every 3 minutes.

The average airway pressure is higher, and a longer filling time is normally available.

To avoid overinflation of the lung, the peaks of pressure during the sigh phase can be limited by the pressure limit P_{max} without impairing the sigh function.

During the sigh phase, the "Volume not constant" alarm is disabled.



SIMV

Synchronised Intermittent Mandatory Ventilation

Combination of machine ventilation and spontaneous breathing.

SIMV enables the patient to breathe spontaneously in regular prescribed intervals between mandatory mechanical ventilation strokes that ensure a minimum ventilation. This minimum ventilation is defined by two set values, tidal volume (V_T) and ventilation frequency (f). The minimum ventilation is the product of $V_T \times f$.

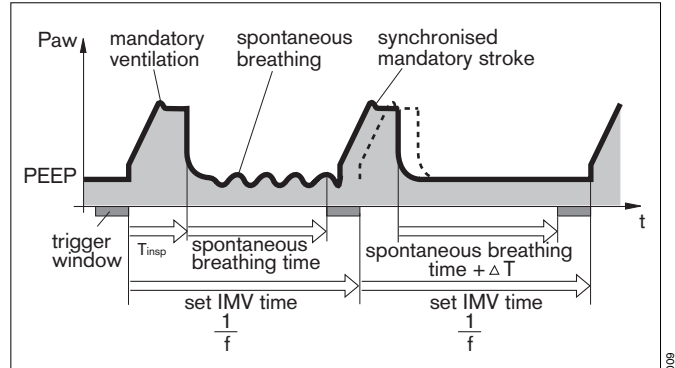
The ventilation pattern is programmed by the following set values: tidal volume V_T , Insp. Flow, frequency f and inspiration time T_{insp} . To prevent the mandatory ventilation stroke being applied during spontaneous expiration, the Flowtrigger of the machine ensures that the ventilation stroke is triggered within a "trigger window" and synchronised with the patient's spontaneous inspiration.

The "trigger window" is 5 seconds long in adult mode and 1.5 seconds long in paediatric mode. If the expiration times are less than 5 seconds or 1.5 seconds respectively, the trigger window covers the entire expiration time.

Since the synchronisation of the mandatory ventilation stroke reduces the effective SIMV time and therefore would normally result in an undesirable increase in the effective SIMV frequency, Evita 2 dura adds back the reduced SIMV time by prolonging the subsequent spontaneous breathing phase by the SIMV time difference ΔT – thus preventing an increase in SIMV frequency. The frequency parameter f remains constant. This parameter, in combination with the tidal volume V_T , sets the minimum ventilation. If the patient has breathed in a considerable inspiratory volume at the beginning of the trigger window, the machine reduces the subsequent mandatory ventilation stroke by shortening the time for the inspiratory flow phase and the inspiration time. In this way, the tidal volume V_T remains constant, and overinflation of the lungs is avoided.

During the spontaneous breathing phases, the patient can be assisted with pressure by ASB pressure support.

During the weaning process, the frequency f /breath rate on the ventilator is gradually reduced, the clinician thereby reduces the mandatory minute volume. The increase in expiratory time allows for a longer spontaneous breathing window, eventually the required minute volume level should be met by the patients' spontaneous breath rate.



ASB

Assisted Spontaneous Breathing

Pressure support for insufficient spontaneous breathing.

The function of the machine in assisting insufficient spontaneous breathing is similar to that of the anaesthetist who manually assists and monitors the patient's spontaneous breathing by feeling the breathing bag.

The machine takes over part of the inhalation function, with the patient maintaining control of spontaneous breathing.

The CPAP system supplies the spontaneously breathing patient with breathing gas, even if the inspiration effort is weak.

The pressure support of the ASB system is started:

- when the spontaneous inspiration flow reaches the set value of the Flowtrigger, or at the latest
- when the spontaneous inspired volume exceeds 25 mL (12 mL in paediatric mode).

The machine then produces an increase in pressure up to the preselected ASB pressure P_{ASB} , which is adjustable to the breathing requirement of the patient.

The time for this pressure increase («Ramp») is adjustable from 0.05 seconds to 2 seconds.

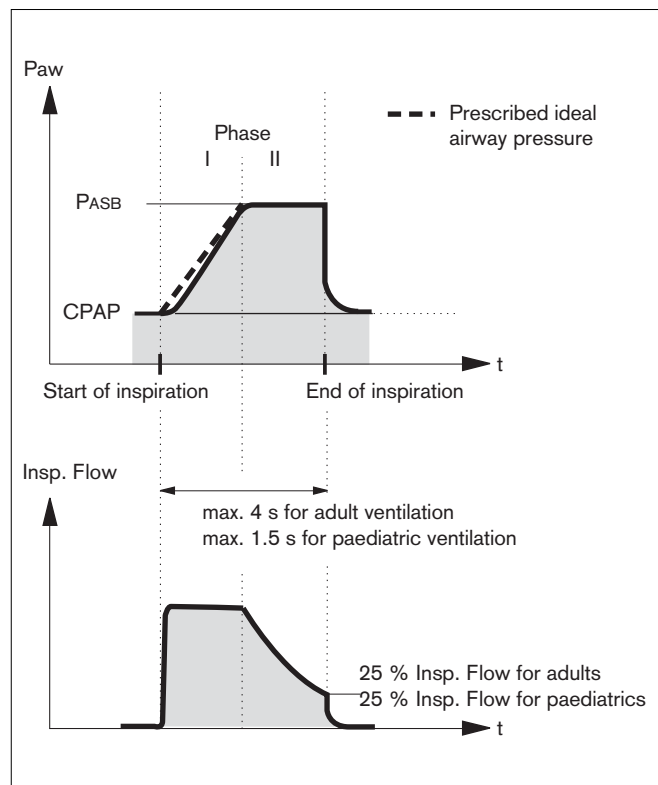
With a rapid increase in pressure Evita 2 dura supports the insufficient spontaneous breathing of the patient with a high peak flow.

With a slow increase in pressure Evita 2 dura begins gently with regular inspiratory flow. The patient has to take over more breathing effort, and so the tone of the breathing muscles gradually improves.

With patient-adjusted pressure increase and the preset ASB level, P_{ASB} , the patient's own breathing activity defines the required inspiration flow, which can rise to 2 L/s in 8 ms.

ASB is terminated:

- when the inspiration flow returns to zero during phase I, i.e. when the patient exhales or fights the ventilator, or
- when the inspiration flow in phase II falls below a certain ratio of the maximum value previously supplied:
25 % Insp. Flow for adults
25 % Insp. Flow for paediatrics
or
- at the latest after 4 seconds (1.5 seconds in paediatric ventilation) if the other two criteria have not come into operation.
If this 4-second criterion occurs three times in succession, Evita 2 dura sounds an alarm and warns of a possible leak in the ventilation system.



BIPAP

Biphasic Positive Airway Pressure

The BIPAP ventilation mode is a pressure/time-cycled ventilation mode in which the patient can always breathe spontaneously. BIPAP is therefore often described as a timed alternation between two CPAP levels.*

The time-cycled change of pressure produces controlled ventilation corresponding to the pressure-controlled ventilation PCV. However, the constant option of spontaneous breathing allows the transition from controlled breathing to independent spontaneous breathing to take place smoothly over the course of the weaning phase, without requiring any change in the ventilation mode. To adapt easily to the patient's spontaneous breathing pattern, the change-over from expiratory pressure level to inspiratory pressure level, and also the change-over from inspiratory pressure level to expiratory pressure level, are synchronised with the patient's spontaneous breathing.

The frequency of the change-over is kept constant, even with patient synchronisation, by defining a trigger time window with a fixed time constant.

The "trigger window" is 5 seconds long in adult mode and 1.5 seconds long in paediatric mode. For expiration times shorter than 5 seconds or 1.5 seconds respectively, the trigger window covers the entire expiration time. At P_{insp} level, the "trigger window is" $1/4 \times T_{insp}$ seconds long.

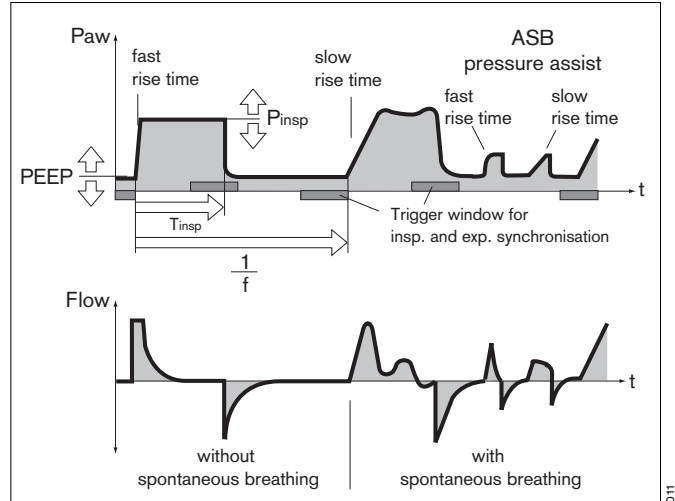
This smooth adaptation to the patient's spontaneous breathing requires less sedation, so that the patient returns to spontaneous breathing more rapidly.

As in all pressure-controlled ventilation modes, the patient is not prescribed a fixed tidal volume (V_T). The tidal volume results principally from the pressure difference between the settings for PEEP and P_{insp} .

The display of the tidal volume measured on expiration, V_{Te} , is used to set the required difference between the two pressure levels. Any increase in this difference will cause an increased BIPAP ventilation stroke.

Changes in lung compliance and airways, as well as active 'fighting' by the patient can lead to changes in tidal volume. This is a desired effect in this ventilation mode.

With the knowledge that the tidal volume, and therefore the minute volume, are not constant, the alarm limits for minute volume must be adjusted with care.



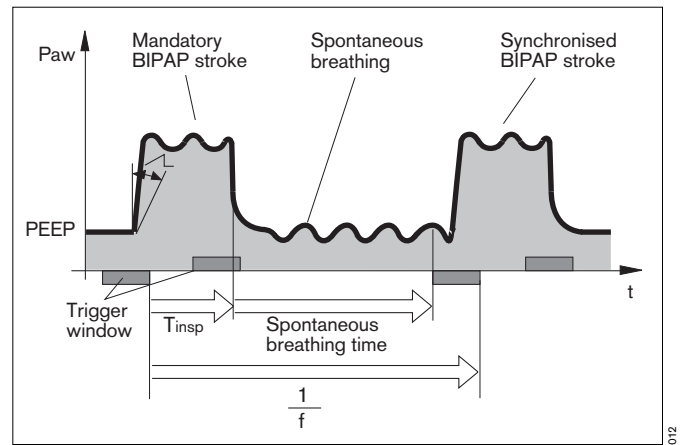
* Bibliography (1), (2), (3), (4), (5), (6), (7), (8), (9), (10) page 131

Using BIPAP

As with IPPV, the time pattern is set using the basic setting parameters of frequency f and inspiration time T_{insp} . The lower pressure level is set with the PEEP parameter, while the upper level is set with P_{insp} . When switching over from IPPV to BIPAP mode – while retaining the time pattern – only the P_{insp} setting needs to be changed.

The steepness of the increase from the lower pressure level to the upper pressure level is controlled by the »Ramp« setting. The effective time for the increase in pressure cannot be greater than the set inspiratory time T_{insp} .

This precaution ensures that the upper pressure level P_{insp} is reached safely during inspiration. The transition from controlled ventilation via the weaning phase to fully spontaneous breathing is achieved by a gradual reduction of the inspiratory pressure P_{insp} and/or frequency f .



BIPAP_{Assist}

Biphasic Positive Airway Pressure Assisted

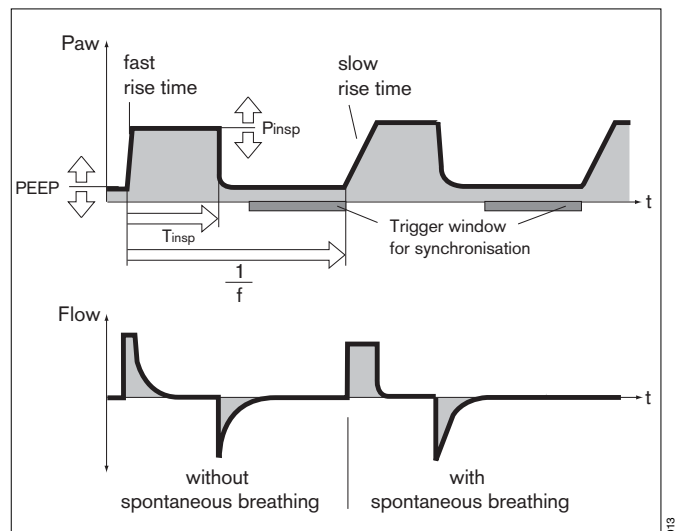
Pressure-controlled, assisted ventilation

The inspiratory strokes are the same as for BIPAP, except that the change from P_{insp} to PEEP is not synchronised with expiration by the patient. The duration of P_{insp} depends on T_{insp} . The patient can breathe spontaneously throughout the ventilation process.

Every detected spontaneous breathing activity by the patient triggers a synchronised inspiration stroke.

A non-synchronised inspiratory stroke is triggered by the machine at the latest upon expiry of the inspiration time defined by » f « and » T_{insp} «.

For all patients, from those unable to breathe spontaneously to those breathing spontaneously before being weaned off the ventilator.



MMV

Mandatory Minute Volume Ventilation

In contrast to SIMV, the MMV ventilation mode gives mandatory breathing only if spontaneous breathing is not yet sufficient and has fallen below a pre-selected minimum ventilation. This minimum ventilation is controlled by the two set values tidal volume V_T and frequency f , and results from the product $V_T \times f$.

Unlike SIMV, the mandatory strokes are not given regularly but only in cases of insufficient ventilation.

The frequency of mandatory strokes is determined by the level of spontaneous breathing: if spontaneous breathing is sufficient, mandatory strokes are not applied at all: If spontaneous breathing is not sufficient, intermittent mandatory strokes of the set tidal volume V_T are applied. If there is no spontaneous breathing at all, the mandatory strokes are applied at the set frequency f .

Evita 2 dura continuously monitors the difference between spontaneous breathing and the set minimum ventilation. As soon as the balance becomes negative, because spontaneous breathing is no longer sufficient, Evita 2 dura applies a mandatory ventilation stroke with the set tidal volume V_T , so that the balance is again positive.

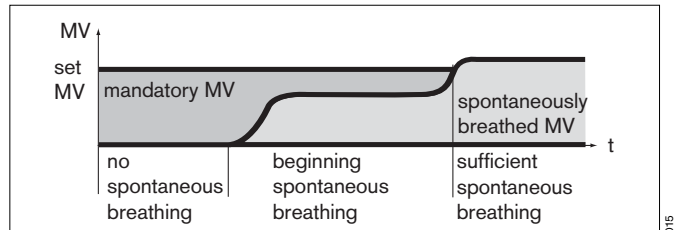
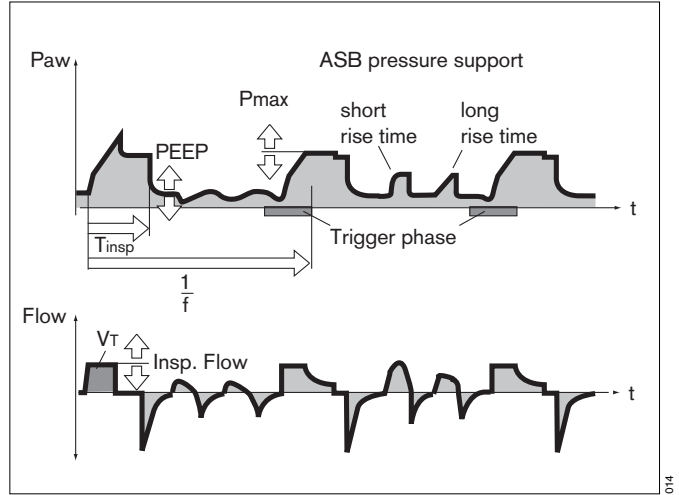
Experience shows that patients breathe very irregularly. Phases of weak breathing alternate with phases of heavy breathing. In order to allow for these individual fluctuations, the balancing process also takes account of the extent by which the set minimum ventilation has been exceeded. This positive allowance is progressively reduced to zero by Evita 2 dura within a maximum of 7.5 seconds after apnoea.

In other words, the response time of Evita 2 dura before activating mandatory ventilation is automatically adapted to the preceding cycle of spontaneous breathing:

If this spontaneous breathing was close to the minimum ventilation, the machine responds rapidly within the cycle time ($1/f$). By contrast, if the patient's spontaneous breathing was much higher than the set minimum ventilation, Evita 2 dura tolerates a longer breathing pause. In extreme cases of sudden apnoea after a phase of heavy breathing, the response time will be 7.5 seconds plus the trigger time, with a minimum of 1 cycle time ($1/f$).

Response times longer than 15 seconds may only occur if the minimum ventilation with a very low IMV frequency f is set to correspondingly low values.

In this case, Evita 2 dura triggers an apnoea alarm that is cancelled again as soon as the mandatory ventilation strokes have been applied. If the cycle time ($1/f$) is set to a longer period than the $T_{Apnoea} \sqrt{f}$ alarm limit, and if there is no spontaneous breathing between the mandatory ventilation strokes, the apnoea alarm will be regularly triggered.



Example:

$f = 3/\text{min} = \text{cycle time } (1/f) = 20 \text{ seconds}$

$T_{\text{Apnoea}} \sqrt{A} = 15 \text{ seconds}$

This system is designed to prevent mandatory ventilation being prematurely triggered in the event of irregular spontaneous breathing, whilst at the same time giving an alarm for any long period of low ventilation.

Measurements

Flow measurement

Regardless of whether ventilation is volume-controlled or pressure-controlled, positive pressures are generated in both the breathing system and patient lung during the inspiration phase. Depending on the ratio of lung compliance to hose system compliance, the volume delivered by the ventilator is distributed to the patient's lung and to the hose system installed between the ventilator and patient. Deviations in the measured expiration flow and derived values, such as the minute volume and breath volume, are low for adult patients, due to their relatively high lung compliance in relation to the much lower compliance of the ventilation hoses.

However, since only the volume attained and surrendered by the lung is relevant to the efficiency of ventilation, and since higher differences are possible during paediatric ventilation, Evita 2 dura provides basic compensation for hose compliance during ventilation.

Compensation of the effect of hose system compliance

During the device check before ventilation, Evita 2 dura determines the compliance of the ventilation hoses, and then, during ventilation, compensates for the effect of compliance on volumetric flow measurement.

Depending on the airway pressure, Evita 2 dura increases the tidal volume by the amount that remains in the ventilation hoses.

In addition to hose system compliance, flow/volume measurement is influenced by the environmental factors of temperature and humidity and by leaks in the hose system. Evita 2 dura takes these factors into account and corrects the settings and measured values accordingly.

Description

Measurements

Conversion according to ambient conditions

The volume occupied by a gas depends on the ambient conditions of temperature, pressure and humidity. In lung physiology, the minute volume and tidal volume are related to the ambient conditions in the lung: 37 °C body temperature, pressure in the lung, 100 % relative humidity.

The flow and volume values measured under these conditions are marked with BTPS*. On the other hand, medical gases from cylinders or from the central supply are dry (approx. 0 % r.h.) and are delivered by the ventilator at 20 °C. The flow and volume values measured under these conditions are marked NTPD**. The difference between measured values under NTPD and BTPS conditions is typically approx. 12 %.

Example: a tidal volume of 500 mL NTPD is increased to 564 mL BTPS by heating to 37 °C and humidifying to 100 % r.h. Evita 2 dura delivers the tidal volume after conversion, so that the set tidal volume is effective in the lung under BTPS conditions.

Automatic leakage compensation

Evita 2 dura determines the difference between the delivered flow on the inspiration side and the measured flow on the expiration side. This difference provides a measure of the amount of leakage and is displayed by Evita 2 dura as the leakage minute volume MV_{leak} . Evita 2 dura can compensate for this leakage in volumecontrolled ventilation.

Example:

Tidal volume setting $V_T = 500$ mL,
10 % leakage in tube.

Leakage compensation Off

Evita 2 dura delivers 500 mL. This is indicated as the inspiratory tidal volume V_{Ti} . 50 mL escape as leakage during inspiration, and 450 mL reach the lung. 450 mL are expired, and 45 mL again escape as leakage. A tidal volume of 405 mL is measured on the expiration side and indicated as V_{Te} . With a ventilation rate of 10 strokes per minute, a minute volume of 5.0 L/min is delivered on the inspiration side and a minute volume of 4.05 L/min is measured on the expiration side. The lung is ventilated with an MV of 4.5 L/min.

Without leakage compensation, the set V_T determines the volume delivered by Evita 2 dura.

* BTPS = Body Temperature, Pressure, Saturated.

** NTPD = Normal Temperature Pressure Dry.

Leakage compensation On

With automatic leakage compensation, Evita 2 dura delivers 550 mL on the basis of the measured leakage minute volume, instead of the 500 mL set. 500 mL enter the lung and the displayed inspiratory tidal volume V_T is 500 mL. The volume of 450 mL measured on the expiration side is displayed without compensation, even when leakage compensation is activated. The minute volume measured on the expiration side is 4.5 L/min and is also uncompensated. If this were not so, the alarm for a low minute volume could be inhibited by the expiratory leakage compensation. Evita 2 dura must always emit an alarm if the minute volume is too low.

With leakage compensation, the set V_T determines the volume to be delivered to the patient.

This example has been simplified:

In fact, the calculated leakage correction takes into account the pressures in the hose system. A higher percentage volume is lost on the inspiration side than on the expiration side because the pressure during inspiration is higher. The displayed leakage minute volume MV_{leak} is based on the mean pressure P_{mean} .

The leakage minute volume MV_{leak} also takes the inspiratory leaks into account. The sum of the minute volume MV + the leakage minute volume MV_{leak} is consequently greater than the inspiratory minute volume delivered to the patient.

Unlimited volume compensation is inappropriate.

Evita 2 dura compensates for losses of up to 100 % of the set tidal volume V_T . Due to technical tolerances, a small leakage minute volume may be displayed even if the hose system is leakproof.

Rapid-Shallow-Breathing RSB

The Rapid Shallow Breathing index (RSB)* is the quotient of the spontaneous breathing frequency (spontaneously breathed breaths per minute) and the tidal volume

$$\text{RSB [1/(min x L)]} = \frac{f_{\text{spn}} [1/\text{min}]}{V_T[\text{L}]}$$

The lower the RSB index for a patient with spontaneous breathing, the more probably he or she can be weaned successfully. The significance of the RSB index is due to the fact that patients who can be weaned successfully tend to have a lower spontaneous breathing frequency and a higher tidal volume than those who are not yet ready to be weaned. In their 1991 study*, Yang and Tobin showed that the RSB index is an effective instrument for predicting the success of an attempt to wean the patient. Patients with an RSB index <100 1/(min x L) were weaned with a probability of 80 %, while 95 % of those with an RSB index > 100 were not yet ready to be weaned. Evita 2 dura indicates the RSB index in CPAP/ASB and PPS modes.

Negative Inspiratory Force NIF

The Negative Inspiratory Force index (NIF)** measures the patient's maximum inhalation effort after exhaling. The patient system is closed during measurement of the NIF. This value is also known as the Maximum Inspiratory Pressure (MIP).

As a result of the inhalation effort during manually extended expiration, the patient generates a negative pressure in relation to PEEP. The probability that the patient can be weaned successfully increases with the magnitude of this negative pressure. Patients with a NIF < -30 mbar can in all probability be weaned successfully, while those with a NIF of up to -20 mbar will most probably prove unsuccessful.

Evita 2 dura determines the NIF value during manually extended expiration. The patient system closes following expiration by the patient while the »Exp. hold« key is held down and Evita 2 dura measures the maximum inhalation effort made by the patient. The NIF is measured as a pressure against PEEP. The measuring procedure is ended when the »Exp. hold« key is released or after not more than 15 seconds. The last measured NIF value and the time of measurement are shown in Table 2 on the screen.

* Bibliography (8), page 131

** Bibliography (9), (10), page 131

Insp. O₂ concentration during medicament nebulisation

Use only medicament nebuliser 84 12 935 (white central body).

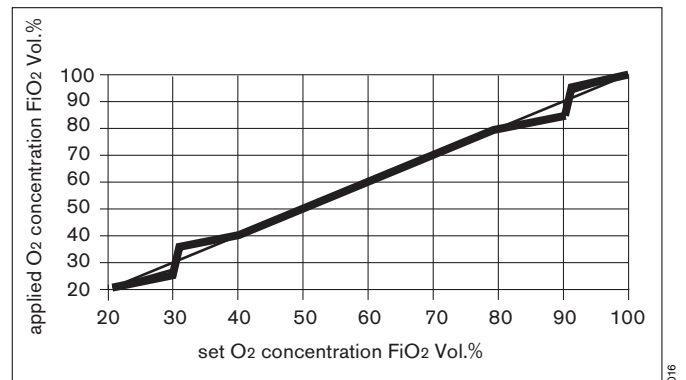
If other medicament nebulisers are used, considerable deviations may occur in the tidal volume and inspiratory O₂ concentration!

To minimise the deviation from the set O₂ concentration, Evita 2 dura generates a mixed gas to drive the medicament nebuliser.

In adult ventilation, this mixed gas is generated by switching over the compressed gases (medical air and oxygen) in synchronisation with inspiration.

In paediatric ventilation, the nebuliser is operated continuously, with medical air or oxygen in alternation. The drive gas of the medicament nebuliser therefore roughly corresponds to the set FiO₂.

The graph shows the possible deviations of the applied O₂ concentration as a function of the set FiO₂ at a minimal inspiratory flow (15 L/min) in adult ventilation, or at ventilation frequencies above 12 bpm in paediatric ventilation.



Abbreviations**Abbreviation Definition**






















APRV	Airway Pressure Release Ventilation Spontaneous breathing at continuous positive airway pressure with short-term pressure release
ASB	Assisted Spontaneous Breathing Pressure supported spontaneous breathing
BIPAP	Biphasic Positive Airway Pressure Ventilation mode for spontaneous breathing at continuous positive airway pressure with two different pressure levels
BIPAP _{Assist}	Biphasic Positive Airway Pressure Assisted Ventilation mode for assisted ventilation with continuous positive airway pressure with two different pressure levels
Body Wt	Body weight (kg)
bpm	breath per minute
BTPS	Body Temperatur, Pressure. Saturated Measured values based on the condition of the patient's lungs, with body temperature 37 °C, steam-saturated gas, atmospheric pressure
C	Compliance
CPAP	Continuous Positive Airway Pressure Breathing with continuous positive pressure in the airways
etCO ₂	End-expiratory CO ₂ concentration
f	Frequency
Fail to cycle	Breathing cycle failure. Machine detects no inspiration
f _{Apnoea}	Frequency setting for apnoea ventilation
FeCO ₂	Expiratory CO ₂ concentration
FiO ₂	Inspiratory O ₂ concentration
Flow	Set value of the maximum inspiratory flow
f _{mand}	Mandatory mechanical portion of overall breathing frequency
f _{spn} , f _{spont}	Spontaneous breathing portion of overall breathing frequency
HME	Heat Moisture Exchanger Heat and moisture exchanger
I : E	Ratio of Inspiration to Expiration
Int. PEEP	Intermittent Positive End-Expiratory Pressure = Sigh
IPPV	Intermittent Positive Pressure Ventilation
IPPV _{Assist}	Trigger Assist Intermittent Positive Pressure Ventilation
IRV	Inversed Ratio Ventilation Ventilation with inversed inspiration/expiration ratio

Abbreviation**Definition**

ISO 5369	International standard for mechanical ventilators – "Lung Ventilation"
MMV	Mandatory Minute Volume Ventilation
MV	Minute Volume
MV _{leak}	Leakage minute volume
MV _{spn}	Spontaneous breathed minute volume
NIF	Negative Inspiratory Force Maximum inhalation effort
O ₂	Set value for inspiratory oxygen concentration [Vol.%]
PASB	Set value of ASB pressure support
P _{aw}	Airway pressure
PEEP	Positive End-Expiratory Pressure
PEEP _i	intrinsic Positive End-Expiratory Pressure
P _{high}	Set value of the upper pressure level APRV
P _{insp}	Set value of the upper pressure level in BIPAP
P _{low}	Value set for lower pressure level in APRV
PLV	Pressure Limited Ventilation
P _{max}	Set value for pressure limited ventilation
P _{mean}	Mean airway pressure
P _{peak}	Peak pressure
P _{plat}	End-inspiratory airway pressure
R	Resistance
Ramp	Setting for the temporary pressure increase in ASB
RSB	Rapid Shallow Breathing Quotient of spontaneous breathing frequency and tidal volume
SIMV	Synchronized Intermittent Mandatory Ventilation
T	Inspiratory breathing gas temperature
T _{Apnoea}	Apnoea alarm time
T _e	Expiration time
TGI	Tracheale Gas Insufflation
T _{high}	Time for the upper pressure level in APRV
T _{insp}	Set value of the inspiratory time
T _{low}	Time for the lower pressure level in APRV
\dot{V} CO ₂	CO ₂ production [L/min]
V _{ds}	Serial dead space
VT	Setting for tidal volume
VT _{Apnoea}	Setting for tidal volume of apnoea ventilation
VTASB	Inspiratory breathing volume during an ASB stroke

Abbreviation	Definition
V_{Te}	Expiratory tidal volume
V_{Ti}	Inspiratory tidal volume
V_{trap}	Volume trapped in the lung by intrinsic PEEP, and exhaled during subsequent expiration.

Symbols

Symbol	Definition
	Switch medicament nebuliser on/off
O ₂ ↑ Suction	Activate/deactivate oxygen enrichment for bronchial suction
Insp. hold	Start manual inspiration
Exp. hold	Manually extend the expiration phase
	Bright/dark screen brightness setting
	Manual printer logging
	Switch help function on/off
Freeze 	"Freeze" curves in screen
	Suppress audible alarm for 2 minutes
Alarm Reset	Acknowledge alarms
	Protect ventilation parameters and ventilation mode
	Standby/Operation
1  2 	Select other combination of measured values
	Select other curve(s)
	Lower/upper alarm limit
	Observe Instructions for Use!
	Protection class Type B
	Protection class Type BF
	Insert flow sensor
	Unlocking expiration valve
	Evita Remote Pad
	Nurse call
Exp.	Expiration port (GAS RETURN)
Insp.	Inspiratory port (GAS OUTPUT)*
	Gas outlet (EXHAUST – NOT FOR SPIROMETER)*
	Earth

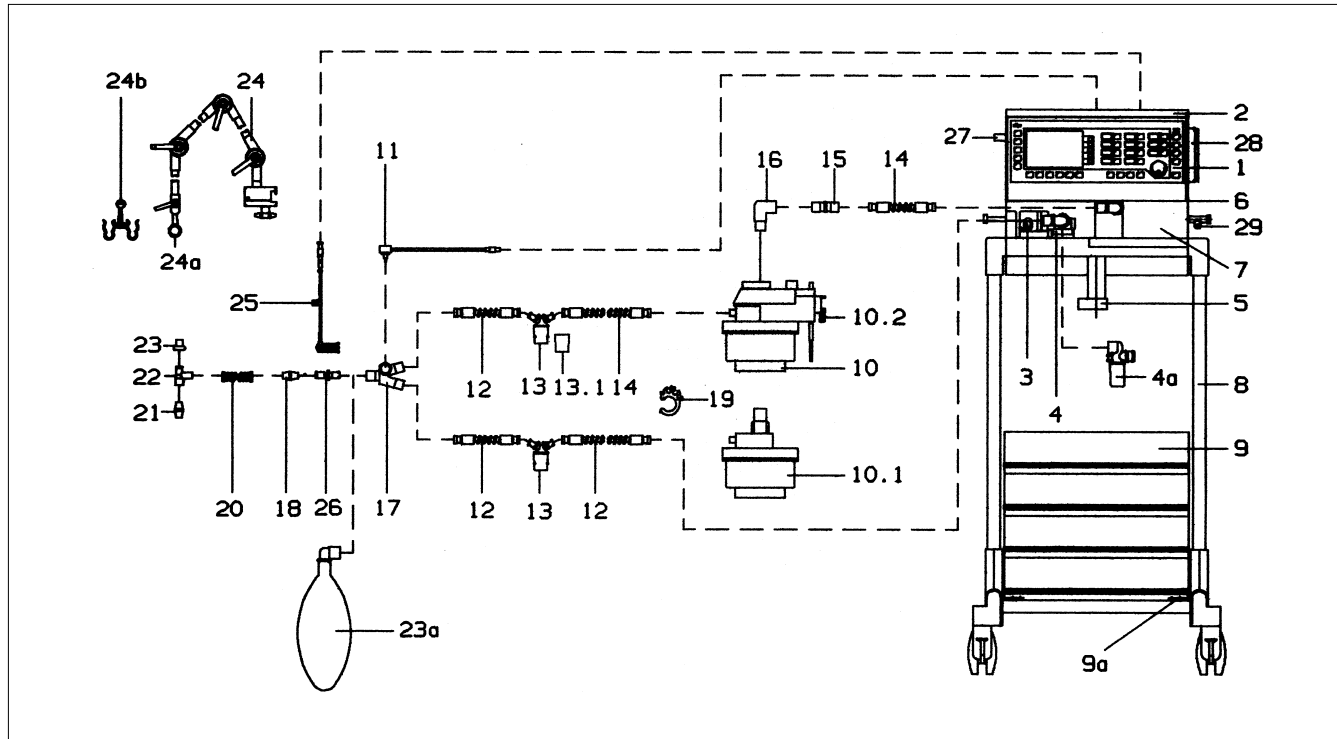
* Additional, depending on equipment configuration

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Parts List

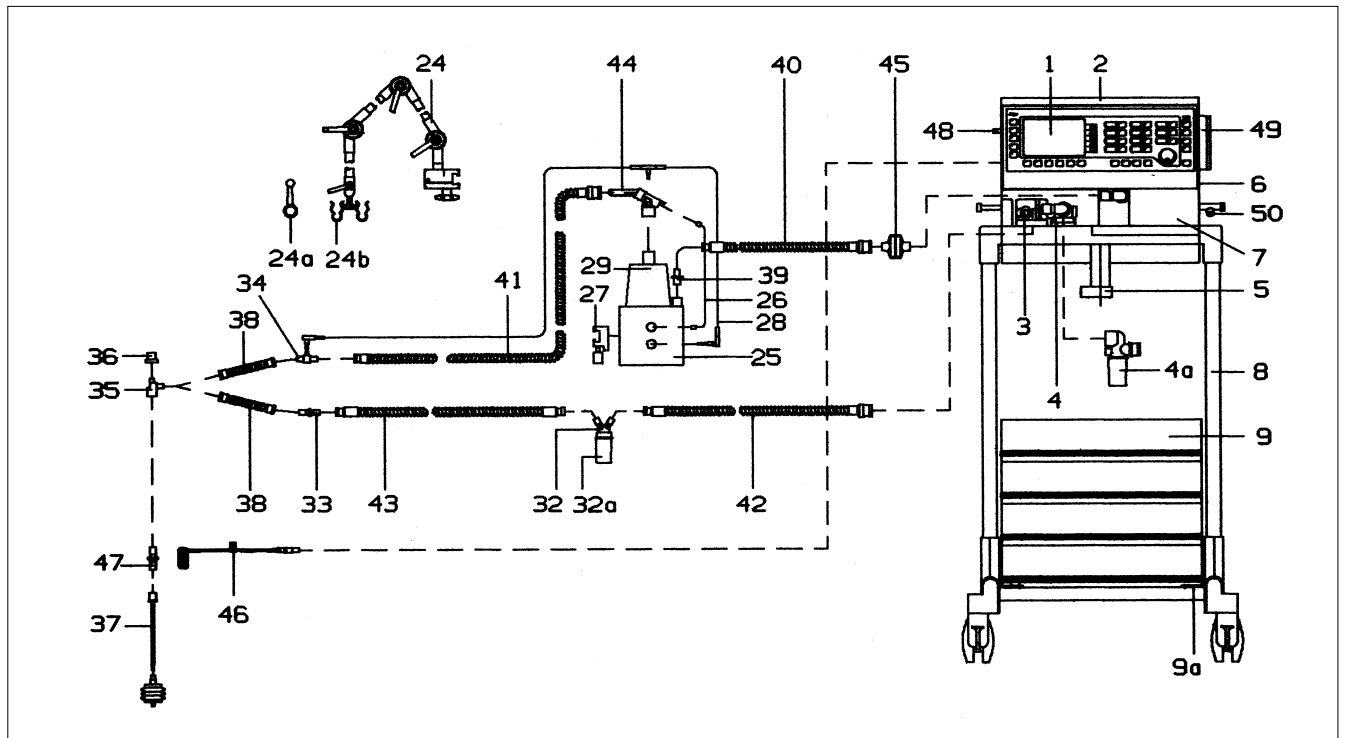
For adult ventilation



Item No.	Name/Description	Order No.
1	Evita 2 dura	84 11 800
2	Instrument tray	84 11 621
3	Flow sensor (set of 5)	84 03 735
4	Expiration valve	84 10 580
5	Bracket (for Aquapor)	84 11 956
6	O ₂ sensor housing	68 50 645
7	Ambient-air filter	84 12 384
7a	Cooling-air filter (back of Evita 2 dura, not illust.)	84 12 384
8	"EvitaMobil" trolley (high)	84 11 950
8a	"EvitaMobil" trolley (low)	84 11 965
8b	Cabinet 8H, 360 mm high (4 drawers)	M 31 796
8c	Cabinet 4H (2 drawers) (not illust.)	M 31 795
8d	"EvitaMobil" cylinder holder set (not illust.)	84 11 970
8e	Breathing air compressor (not illust.)	84 13 890
10	Aquapor (220 to 240 V)	84 05 020
	Aquapor (110 V)	84 05 199
10.1	Patient part, Aquapor	84 05 029
10.2	Set of spare brackets	84 03 345
11	Temperature sensor	84 05 371
12-23	Hose set, adult (blue socket)	84 12 092
12	Spiral hose, adult, silicone 0.6 m	21 65 627
13-13.1	Water trap	84 04 985

Item No.	Name/Description	Order No.
13.1	Water container	84 03 976
14	Spiral hose, adult, silicone 0.35 m	21 65 619
15	Connector	M 25 647
16	ISO elbow connector	M 25 649
17	Y-piece, straight	84 05 435
18	Catheter connector, straight, size 12.5 (set of 10)	M 23 841
19	Hose clamp	84 03 566
20	Corrugated hose 0.32 m	84 02 041
21	Catheter connector, adult	
	Set of catheter connectors, adult	
	Sizes 6 to 12 (set of 12)	84 03 685
22	Adaptor, adult	84 03 076
23	Cap (set of 5)	84 02 918
23a	Adult test lung (bag)	84 03 201
24-24b	Hinged arm or	84 09 609
	Quick-fix hinged arm 2	2M 85 706
24a	Bracket	84 09 746
24b	Hose clamp	84 09 841
25	CO ₂ main flow sensor	68 70 300
26	Cuvette, adult	68 70 279
27	Holder for parking CO ₂ sensor	84 12 840
28	Short GA holder set	84 11 615
29	Y-piece parking rest	84 11 784

For paediatric ventilation



Item No.	Name/Description	Order No.	Item No.	Name/Description	Order No.
1	Evita 2 dura	84 11 800	31	Single-strand wire 1.5 m (not illustr.)	84 11 050
2	Instrument tray	84 11 621	32-43	Hose set, paediatrics (Fisher & Paykel)	84 12 081
3	Flow sensor (set of 5)	84 03 735	32-32a	Condensation trap, expiration	84 09 627
4	Expiration valve (expiration valve)	84 10 580	32	Water container	84 03 976
5	Bracket (for Aquapor)	84 11 956	33	Double conical connector	84 09 897
6	O ₂ sensor housing	68 50 645	34	Temperature sensor mounting	84 11 044
7	Ambient-air filter	84 12 384	35	Adapter K90	84 03 075
7a	Cooling-air filter (back of Evita 2 dura, not illust.)	84 12 384	36	Cap 5x	84 01 645
8	"EvitaMobil" trolley (high)	84 11 950	37	Bellows, paediatric, complete	84 09 742
8a	"EvitaMobil" trolley (low)	84 11 965	38	Corrugated hose, flex., 0.13 m	84 09 634
8b	Cabinet 8H, 360 mm high (4 drawers)	M 31 796	39	Catheter connector, size 11	M 19 351
8c	Cabinet 4H (2 drawers) (not illust.)	M 31 795	40	Spiral hose, paediatric, silicone 22/10, 0.40 m	21 65 856
8d	"EvitaMobil" cylinder holder set (not illust.)	84 11 970	41	Spiral hose, paediatric, silicone 22/10, 1.10 m	21 65 651
8e	Breathing air compressor (not illust.)	84 13 890	42	Spiral hose, paediatric, silicone 22/10, 0.60 m	21 65 821
24-24b	Hinged arm or Quick-fix hinged arm 2	84 09 609 2M 85 706	43	Spiral hose, paediatric, silicone 10/10, 0.60 m	21 65 848
24a	Bracket	84 09 746	44	Hose heater 1.10 m	84 11 045
24b	Hose clamp	84 09 841	45	Bacterial filter	MX 02 650
26-28	Humidifier, basic unit MR 730 (Fisher & Paykel)	84 11 046	46	CO ₂ main flow sensor	68 70 300
26	Hose heater adapter	84 11 097	47	Cuvette, paediatrics	68 70 280
27	Mounting set (clamps for rail)	84 11 074	48	Holder for parking CO ₂ sensor	84 12 840
28	Double temperature sensor	84 11 048	49	Short GA holder set	84 11 615
29-30	Humidifier chamber MR 340	84 11 047	50	Y-piece parking rest	84 11 784
30	Filter paper (set of 100, not illustr.)	84 11 073			

Order List

Name/Description	Order No.	Name/Description	Order No.
Basic unit		Special accessories	
Evita 2 dura	84 11 800	Instrument tray	84 11 621
Accessories required for operation		Holder for quick reference manual	84 11 615
Hinged arm	84 09 609	Wall bracket, module 2000 Type 13 alternative to trolley	84 08 613
or		Pneumatic medicament nebuliser	84 12 935
Quick-fix hinged arm 2	2M 85 706	Flow sensor cover	84 14 714
O2 connecting hose 3 m, blue/white	M 29 231	For manual ventilation:	
or		Resutator 2000	21 20 046
O2 connecting hose 5 m, blue/white	M 29 251	Paediatric Resutator 2000	21 20 984
O2 connecting hose 3 m, neutral colour	M 34 402	Baby-Resutator	21 20 941
or		Hook for Resutator	M 26 349
O2 connecting hose 5 m, neutral colour/white	M 34 403	Adult test lung	84 03 201
Medical air connecting hose 3 m, yellow/black	M 29 239	consisting of:	
or		Mask manifold	M 25 649
Medical air connecting hose 5 m, yellow/black	M 29 259	Catheter connector ISO size 7	M 25 591
Medical air connecting hose 3 m, neutral colour	M 34 408	Breathing bag, 2 L	21 65 694
or		For trolley:	
Medical air connecting hose 5 m, neutral colour	M 34 409	Cabinet unit 8H, 360 mm high	M 31 796
EvitaMobil trolley	84 11 950	For supplying Evita 2 dura with medical air:	
For adult ventilation		Breathing air compressor	84 13 890
Temperature sensor	84 05 371	Standby option	84 13 939
Aquapor EL humidifier	84 14 698	Special voltage transformer	84 13 936
Set of spare brackets	84 03 345	MEDIBUS cable	83 06 488
Hose set, adult	84 12 092	Printer cable	83 06 489
consisting of: patient hoses, water traps, Y-piece, catheter connectors		Options	
For paediatric ventilation		Installation set, Ventilation Plus	84 13 540
Humidifier, basic unit MR 730	84 11 046	Installation set, Monitoring Plus	84 13 545
(Fisher & Paykel)		Installation set, Service Plus	84 13 550
Mounting set (rail brackets)	84 11 074	Installation set, Evita Link	84 11 735
Humidifier chamber MR 340	84 11 047	Installation set, Evita Sat	84 13 035
Double temperature sensor	84 11 048	Modification set – DC module	84 13 034
Single-strand wire, 1.5 m	84 11 050	Modification set – Evita DC power supply MB	84 15 581
Hose set, paediatric (Fisher & Paykel)	84 12 081	Installation set, Capno Plus	84 13 780
consisting of:		Accessories for CapnoPlus:	
hose heater, patient hoses, water traps, Y-piece, catheter connectors	23 47 020	Adult cuvette	68 70 279
Bacterial filter	MX 02 650	Paediatric cuvette	68 70 280
		CO2 main flow sensor	68 70 300
		Bracket for parking CO2 sensor	84 12 840
		Modification set – Mask ventilation (NIV)	84 14 474

Name/Description	Order No.	Name/Description	Order No.
Modification set – Nurse call	84 14 476	Aquapor bowl	84 04 739
Plug for connecting the nurse call	18 46 248	Aquapor float	84 04 738
Modification set – EvitaRemote	84 14 472	Spiral hose, adult, silicone 0.6 m	21 65 627
Upgrade modification set	84 15 794	Spiral hose, adult, silicone 0.35 m	21 65 619
Software 4.14 Evita 2 dura		Water trap	84 04 985
Upgrade modification set	84 14 470	Water container	84 03 976
Software 4.0 plus Evita 2 dura		Hose clamp	84 03 566
Modification set – Nurse call/ 2nd pressure sensor	84 15 570	Connector	M 25 647
		Y-piece	84 05 435
For CO₂ measurement (option)		Catheter connector, straight, size 12.5 (set of 10)	M 23 841
Test filter	68 70 281	Corrugated hose	84 02 041
Calibration set	84 12 710	Adaptor, adult	84 03 076
Test gas cylinder 5 Vol.% CO ₂ , 95 Vol.% N ₂	68 50 435	Set of catheter connectors, adult	84 03 685
		Set of caps (set of 5)	84 02 918
Spare set for sterilisation		ISO elbow connector	M 25 649
Expiration valve (expiration valve)	84 10 580		
		For paediatric ventilation:	
For adult ventilation:		Spiral hose, paediatric, silicone, 22/10, 1.10 m	21 65 651
Hose set, adult	84 12 092	Spiral hose, paediatric, silicone, 22/10, 0.60 m	21 65 821
Patient part for Aquapor	84 05 029	Spiral hose, paediatric, silicone, 10/10, 0.60 m	21 65 848
Temperature sensor	84 05 371	Spiral hose, paediatric, silicone, 22/10, 0.40 m	21 65 856
Pneumatic medicament nebuliser	84 12 935	Corrugated hose flex., 0.13 m	84 09 634
Cuvette, adult	68 70 279	Catheter connectors, size 11 (set of 10)	M 19 351
		Cap	84 01 645
For paediatric ventilation:		Adaptor, paediatric, 90°	84 03 075
Hose set, paediatric (Fisher & Paykel)	84 12 081	Double conical connector	84 09 897
Humidifier chamber MR 340	84 11 047	Temperature sensor mounting	84 11 044
incl. filter paper for humidifier chamber (set of 100)		Condensation trap, expiration	84 09 727
Cuvette, paediatric	68 70 280	Water container	84 03 976
		Hose heater 1.10 m	84 11 045
Replacement parts		Double temperature sensor	84 11 048
For Evita 2 dura:		Adaptor for hose heater	84 11 097
O ₂ sensor capsule	68 50 645	Single-strand wire, 1.5 m	84 11 050
Flow sensor (set of 5)	84 03 735	Humidifier chamber, MR 340, incl. filter paper (set of 100)	84 11 047
Cooling air filter, blue	84 12 384	Filter paper for humidifier chamber (set of 100)	84 11 073
Cooling air filters DC power supply MB	84 15 572	Bacterial filter	MX 02 650
Lithium battery for data protection	18 35 343		
For hinged arm:			
Holder	84 09 746		
Hose clamp	84 09 841		
For adult ventilation:			
Temperature sensor	84 05 371		
Replacement set of lids for Aquapor	84 06 135		
		Technical documentation available on request	

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These Instructions for Use apply only to
Evita 2 dura
with Serial No.:

If no Serial No. has been filled in by Dräger
these Instructions for Use are provided
for general information only and are not
intended for use with any specific machine
or device.



Directive 93/42/EEC
concerning Medical Devices

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