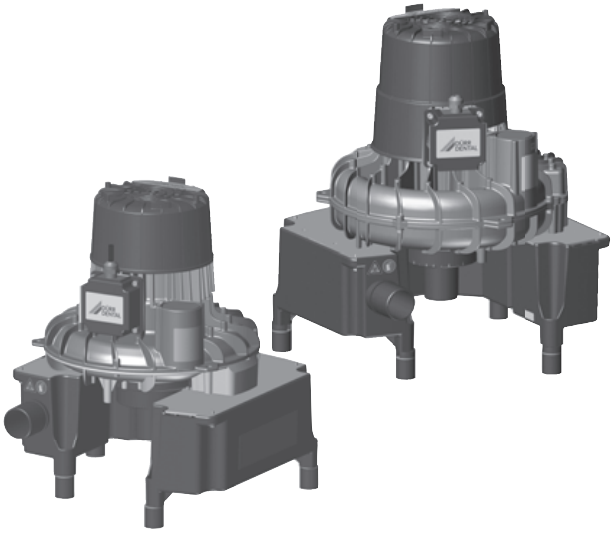
V 600, V 900 S, V 1200 S



EN Installation and operating instructions

CE 0297

DÜRR DENTAL

71311000006L02 2405V003

The current version of the installation and operating instructions is available in the Download Center:

http://qr.duerrdental.com/7131100006

Contents

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Important information

1 About this document

These installation and operating instructions represent part of the unit.

The manufacturer and the distributor will not offer any guarantee or accept any liability for the safe operation and the safe functioning of the unit if the instructions and information in these installation and operating instructions are not complied with.

The German version of the installation and operating instructions is the original manual. All other languages are translations of the original manual. These installation and operating instructions apply to:

V 600
REF: 7127-01; 7127-01/002; 7127-01/021; 7127-02; 7127-02/002

V 900 S
REF: 7131-01; 7131-01/002; 7131-02; 7131-02/002; 7131-02/021

V 1200 S
REF: 7136-02; 7136-02/002; 7136-02/021; 7136-02/003

1.1 Warnings and symbols

Warnings

The warnings in this document are intended to draw your attention to possible injury to persons or damage to machinery.
The following warning symbols are used:


General warning symbol


Warning – dangerous high voltage


Warning – hot surfaces


Warning - automatic start-up of the unit


Biohazard warning

The warnings are structured as follows:

SIGNAL WORD

Description of the type and source of danger

Here you will find the possible consequences of ignoring the warning

› Follow these measures to avoid the danger.

The signal word differentiates between four levels of danger:

– **DANGER**
Immediate danger of severe injury or death
– **WARNING**
Possible danger of severe injury or death
– **CAUTION**
Risk of minor injuries
– **NOTICE**
Risk of extensive material/property damage

Other symbols

These symbols are used in the document and on or in the unit:

Note, e.g. specific instructions regarding efficient and cost-effective use of the unit.


Refer to Operating Instructions.


Wear protective gloves.


Wear protective goggles.


Disconnect all power from the unit.


Refer to the accompanying electronic documents.

Lower and upper temperature limits

Lower and upper humidity limits


Protective ground connection

CE labelling with the number of the notified body

SN Serial number

REF Order number

MD Medical device

HIBC Health Industry Bar Code (HIBC)

 Manufacturer

1.2 Copyright information

2 Safety

The unit has been developed and designed in such a way that dangers are effectively ruled out if used in accordance with the Intended Use. Despite this, the following residual risks can remain:

– Personal injury due to incorrect use/misuse
– Personal injury due to mechanical effects
– Personal injury due to electrical shock
– Personal injury due to radiation
– Personal injury due to fire
– Personal injury due to thermal effects on skin
– Personal injury due to lack of hygiene, e.g. infection

2.1 Intended purpose

The suction unit provides the dental treatment unit with vacuum and volume flow.

2.2 Intended use

Working in combination with the suction unit with treatment unit, suction handpiece and cannula, the media used in dental treatment (e.g. water, saliva, dentine and amalgam) are removed by suction for disposal.
This unit is technically suitable for the aspiration of nitrous oxide (laughing gas). However, when assembling a system for aspiration of nitrous oxide, it is important to ensure that the other components in the system are also suitable for this purpose. Those responsible for setting up the system must assess this and approve and release the system for the aspiration of nitrous oxide.

Operation with nitrous oxide is only permitted if the exhaust air is transported from the unit to the outside of the building.

2.3 Improper use

Any use of this appliance / these appliances above and beyond that described in the Installation and Operating Instructions is deemed to be incorrect usage. The manufacturer cannot be held liable for any damage resulting from incorrect usage. The operator will be held liable and bears all risks.

- Do not use this device to aspirate flammable or explosive mixtures.
- Do not use the unit as a vacuum cleaner.
- Do not use chemicals containing chlorine or foaming chemicals.
- Operation in operating theatres of explosive areas is not permissible.
- The suction unit must not be set up in the immediate surroundings of the patient (minimum distance: 1.5 m).

2.4 General safety information

- Always comply with the specifications of all guidelines, laws, and other rules and regulations applicable at the site of operation for the operation of this unit.
- Check the function and condition of the unit prior to every use.
- Do not convert or modify the unit.
- Comply with the specifications of the Installation and Operating Instructions.
- The Installation and Operating Instructions must be accessible to all operators of the unit at all times.

2.5 Combining devices safely

Take care when connecting units together or to parts of other systems as there is always an element of risk (e.g. due to leakage currents).

- Only connect units when there can be no question of danger to operator or to patient.
- Only connect units when it is safe to do so and when there is no risk of damage or harm to the surroundings.
- If it is not completely clear from the data sheet of the unit that such connections can be safely made or if you are in any doubt, always get a suitably qualified person (e.g. the relevant manufacturer) to verify that the setup is safe.

Where applicable, the requirements for medical products have been taken into account in the development and construction of the device. As a result, this device is suitable for installation within medical supply equipment.

- Where this device is integrated in other medical supply equipment, the requirements of European Union Medical Device Regulation 2017/745 and the relevant standards must be observed.

2.6 Specialist personnel

Operation

Unit operating personnel must ensure safe and correct handling based on their training and knowledge.

- Instruct or have every operator instructed in handling the unit.

The following groups are not permitted to operate or use a commercially operated unit:

- People without the necessary experience and knowledge
- People with reduced physical, sensory or mental capabilities
- Children

Installation and repairs

- Have the manufacturer or a qualified company authorised by the manufacturer perform mounting, new installations, modifications, expansions and repairs.

2.7 Notification requirement of serious incidents

The operator/patient is required to report any serious incident that occurs in connection with the device to the manufacturer and to the competent authority of the Member State in which the operator and/or patient is established/resident.

2.8 Electrical safety

- Comply with all the relevant electrical safety regulations when working on the unit.
- Never touch the patient and unshielded plug connections on the unit at the same time.
- Replace any damaged cables or plugs immediately.

2.9 Only use original parts

- Only use accessories and optional articles named or authorised by the manufacturer.
- Only use only original wear parts and replacement parts.

The manufacturer and distributor accept no liability for damages or injury resulting from the use of non-approved accessories, optional accessories, or from the use of non-original wear parts or replacement parts.

The use of non-approved accessories, optional accessories or non-genuine wear parts / replacement parts (e.g. mains cables) can have a negative effect in terms of electrical safety and EMC.

2.10 Transport

The original packaging provides optimum protection for the unit during transportation.
If required, the original packaging for the unit can be ordered.

The manufacturer and the distributor do not accept liability, even during the warranty period, for damage during transportation due to improper packaging.

– Only transport the unit in its original packaging.
– Keep the packing materials out of the reach of children.

2.11 Disposal

The unit may be contaminated. Instruct the company disposing of the waste to take the relevant safety precautions.

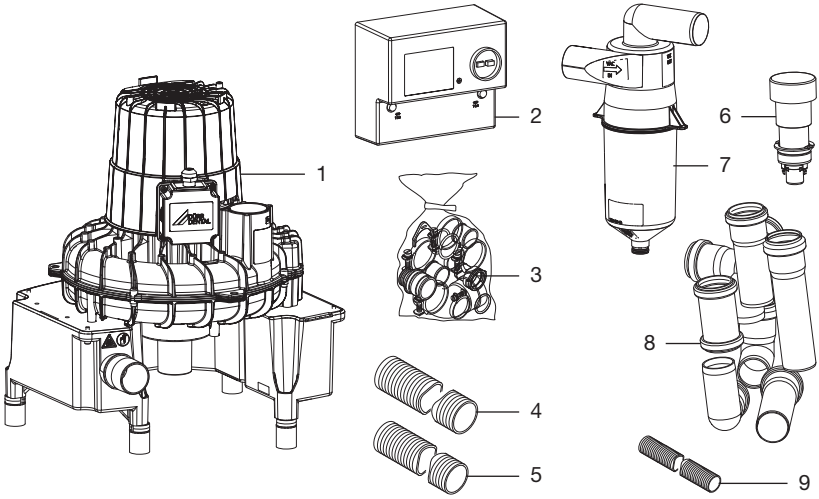
– Decontaminate potentially contaminated parts before disposing of them.
– Uncontaminated parts (e.g. electronics, plastic and metal parts etc.) should be disposed of in accordance with the local waste disposal regulations.
– If you have any questions about the correct disposal of parts, please contact your dental trade supplier.

An overview of the waste keys for Dürr Dental products can be found in the download area:

http://qr.duerrdental.com/P007100155

Product description

3 Overview



1 Suction unit
2 Control box
3 Set of connection fittings
4 Suction hose
5 Exhaust air hose
6 Auxiliary air valve
7 Condensation separator
8 Connection pipes
9 Waste water hose LW 20

3.1 Scope of delivery

The following items are included in the scope of delivery (possible variant-specific deviations due to country-specific requirements and/or import regulations):

V 600, 230 V, 1~, 50 Hz
(basic unit) 7127-01
V 600, 230 V, 1~, 50 Hz
(with accessories) 7127-01/002
V 600, 230 V, 1~, 50 Hz
(Power Tower) 7127-01/021
V 600, 230/400 V, 3~, 50/60 Hz
(basic unit) 7127-02
V 600, 400 V, 3~, 50/60 Hz
(with accessories) 7127-02/002

– Control box
  0700-500-50 for model 7127-01/002
  0732-100-52 for Modell 7127-02/002
– Set of connection fittings
– Suction hose LW 50
– Exhaust air hose LW 50 (aluminium)
– Waste water hose LW 20
– Condensation separator

V 900 S, 230 V, 1~, 50 Hz
(basic unit) 7131-01
V 900 S, 230 V, 1~, 50 Hz
(with accessories) 7131-01/002
V 900 S, 230/400 V, 3~, 50 Hz
(basic unit) 7131-02
V 900 S, 400 V, 3~, 50 Hz
(with accessories) 7131-02/002
V 900 S, 400 V, 3~, 50 Hz
(Power Tower) 7131-02/021

– Control box
  0732-100-50 for model 7131-01/002
  0732-100-52 for Modell 7131-02/002
– Set of connection fittings
– Suction hose LW 50
– Exhaust air hose LW 50 (aluminium)
– Waste water hose LW 20
– Condensation separator

V 1200 S, 230/400 V, 3~, 50 Hz
(basic unit) 7136-02
V 1200 S, 400 V, 3~, 50 Hz
(with accessories) 7136-02/002
V 1200 S, 400 V, 3~, 50 Hz
(Power Tower) 7136-02/021
V 1200 S, 230 V, 3~, 50 Hz
(with accessories) 7136-02/003

– Control box
  0732-100-54 for Modell 7136-02/002
  0732-100-53 for Modell 7136-02/003
– Set of connection fittings
– Suction hose LW 50
– Exhaust air hose LW 50 (aluminium)
– Waste water hose LW 20
– Condensation separator

3.2 Optional items

The following optional items can be used with the device:

Wall bracket 7130-190-00
Console for floor-mounted installation 7130-191-00
Noise reduction hood 7131-991-00
Bacteria filter 0705-991-50
Noise reduction for exhaust air . . . 0730-991-00
Ventilation kit for cabinet installation 7122-981-50
Exhaust air valve 0732-020-00
Bacteria filter with housing * 7120100000

* Only required for V/VS 600, 2x

3.3 Consumables

The following materials are consumed during operation of the device and must be ordered separately:

Orotol plus (2.5 litre bottle) CDS110P6150
MD 555 cleaner (2.5 litre bottle) . CCS555C6150

3.4 Wear parts and replacement parts

The following working parts need to be changed at regular intervals (refer to the "Maintenance" section):

Information about replacement parts is available from the portal for authorised specialist dealers at:
www.duerrdental.net

4 Technical data

4.1 V 600

Electrical data		7127-01
Rated voltage	V	230, 1~
Mains frequency	Hz	50
Active power	kW	1
Nominal current	A	≤ 5.0
Start-up current	A	~ 22
Motor protection *	A	-
Type of protection		IP 24
Protection class		I

* Motor winding overheat protector

Connections		
Suction connection (outside)	mm	∅ 50
Exhaust air connection (external)	mm	∅ 50

Classification in accordance with EN ISO 10637		
Classification based on separation of solids and liquids		Dry suction system
Classification based on flow rate		Type 1

General data		
Max. number of users		2
Duty cycle	%	100 (S1)
Heat generation rate	MJ/h	3.6
Dimensions (H x W x D)	cm	48 x 41 x 39
Weight, approx.	kg	21
Noise level * ca. without housing with housing	 dB(A) dB(A)	 65 57

* Noise level in accordance with ISO 3746

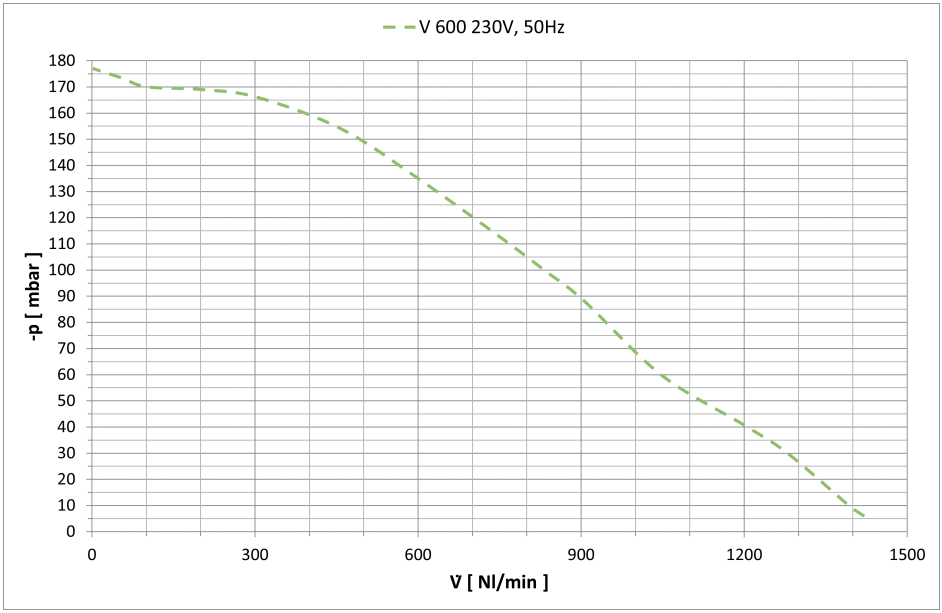
Ambient conditions during storage and transport		
Temperature	°C	-10 - +60
Relative humidity	%	< 95

Ambient conditions during operation		
Temperature	°C	+10 - +40
Relative humidity	%	< 70

Classification

Medical Device Class (MDR)	IIa

Characteristic curve in accordance with EN ISO 10637



Characteristic curve for: 7127-01

4.2 V 600

Electrical data		7127-02	
Rated voltage	V	230, 3~	
Mains frequency	Hz	50	60
Active power	kW	1	1.4
Nominal current	A	≤ 3.8	≤ 4.0
Start-up current	A	~ 16	~ 14
Motor protection *	A	4.5	4.5
Type of protection		IP 24	
Protection class		I	

* Max. permitted setting

Connections		
Suction connection (outside)	mm	∅ 50
Exhaust air connection (external)	mm	∅ 50

Classification in accordance with EN ISO 10637	
Classification based on separation of solids and liquids	Dry suction system
Classification based on flow rate	Type 1

General data			
Max. number of users		2	2
Duty cycle	%	100 (S1)	
Heat generation rate	MJ/h	3.6	5
Dimensions (H x W x D)	cm	48 x 41 x 39	
Weight, approx.	kg	21	
Noise level * ca. without housing with housing	 dB(A) dB(A)	 65 57	 68 60

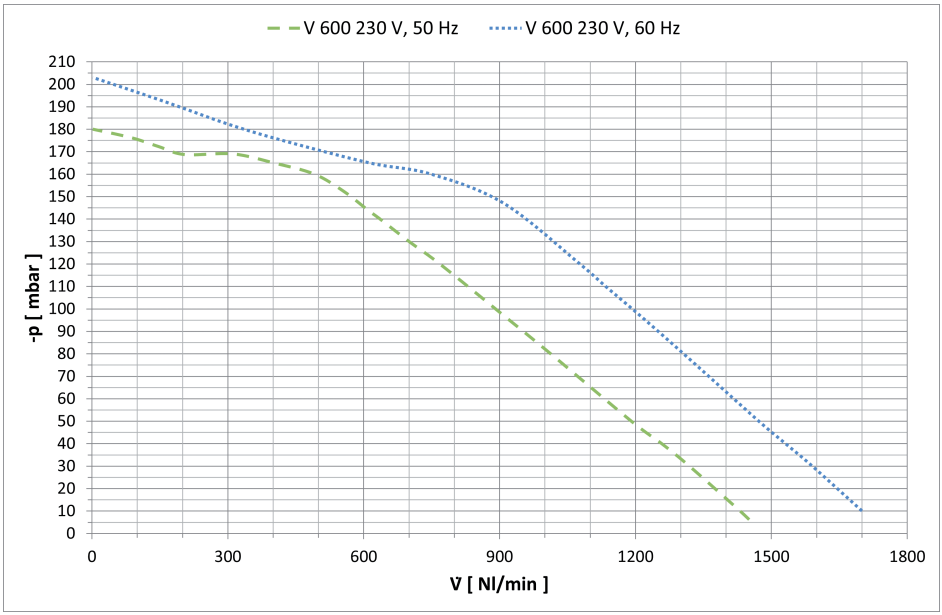
* Noise level in accordance with ISO 3746

Ambient conditions during storage and transport		
Temperature	°C	-10 - +60
Relative humidity	%	< 95

Ambient conditions during operation		
Temperature	°C	+10 - +40
Relative humidity	%	< 70

Classification	
Medical Device Class (MDR)	IIa

Characteristic curve in accordance with EN ISO 10637



Characteristic curve for: 7127-02

4.3 V 600

Electrical data		7127-02	
Rated voltage	V	400, 3~	
Mains frequency	Hz	50	60
Active power	kW	1	1.4
Nominal current	A	≤ 2.2	≤ 2.3
Start-up current	A	~ 16	~ 14
Motor protection *	A	2.5	2.5
Type of protection		IP 24	
Protection class		I	

* Max. permitted setting

Connections		
Suction connection (outside)	mm	∅ 50
Exhaust air connection (external)	mm	∅ 50

Classification in accordance with EN ISO 10637	
Classification based on separation of solids and liquids	Dry suction system
Classification based on flow rate	Type 1

General data			
Max. number of users		2	2
Duty cycle	%	100 (S1)	
Heat generation rate	MJ/h	3.6	5
Dimensions (H x W x D)	cm	48 x 41 x 39	
Weight, approx.	kg	21	
Noise level * ca. without housing with housing	 dB(A) dB(A)	 65 57	 68 60

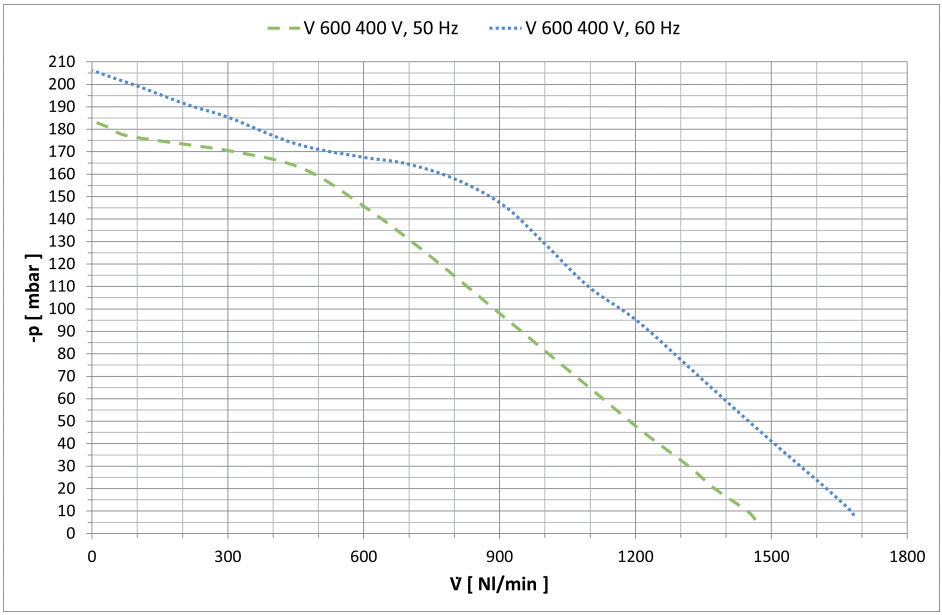
* Noise level in accordance with ISO 3746

Ambient conditions during storage and transport		
Temperature	°C	-10 - +60
Relative humidity	%	< 95

Ambient conditions during operation		
Temperature	°C	+10 - +40
Relative humidity	%	< 70

Classification	
Medical Device Class (MDR)	IIa

Characteristic curve in accordance with EN ISO 10637



Characteristic curve for: 7127-02

4.4 V 900 S

Electrical data		7131-01
Rated voltage	V	230, 1~
Mains frequency	Hz	50
Active power	kW	1.6
Nominal current	A	≤ 7.4
Start-up current	A	~ 32
Motor protection *	A	10.0
Type of protection		IP 24
Protection class		I

* Max. permitted setting

Connections		
Suction connection (outside)	mm	∅ 50
Exhaust air connection (external)	mm	∅ 50

Classification in accordance with EN ISO 10637		
Classification based on separation of solids and liquids		Dry suction system
Classification based on flow rate		Type 1

General data		
Max. number of users		3
Duty cycle	%	100 (S1)
Heat generation rate	MJ/h	5.8
Dimensions (H x W x D)	cm	52 x 41 x 43
Weight, approx.	kg	28
Noise level * ca. without housing with housing	 dB(A) dB(A)	 65 61

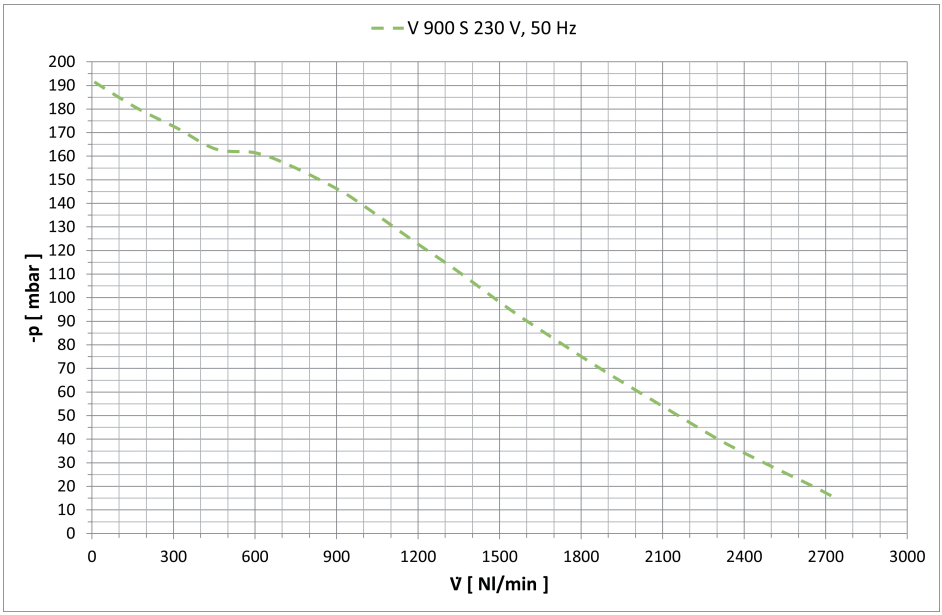
* Noise level in accordance with ISO 3746

Ambient conditions during storage and transport		
Temperature	°C	-10 - +60
Relative humidity	%	< 95

Ambient conditions during operation		
Temperature	°C	+10 - +40
Relative humidity	%	< 70

Classification	
Medical Device Class (MDR)	IIa

Characteristic curve in accordance with EN ISO 10637



Characteristic curve for: 7131-01

4.5 V 900 S

Electrical data		7131-02	
Rated voltage	V	230, 3~	400, 3~
Mains frequency	Hz	50	
Active power	kW	1.7	1.7
Nominal current	A	≤ 6.3	≤ 3.6
Start-up current	A	~ 42	~ 25
Motor protection *	A	6.3	4.0
Type of protection		IP 24	
Protection class		I	

* Max. permitted setting

Connections		
Suction connection (outside)	mm	∅ 50
Exhaust air connection (external)	mm	∅ 50

Classification in accordance with EN ISO 10637		
Classification based on separation of solids and liquids		Dry suction system
Classification based on flow rate		Type 1

General data		
Max. number of users		3
Duty cycle	%	100 (S1)
Heat generation rate	MJ/h	6.1
Dimensions (H x W x D)	cm	52 x 41 x 43
Weight, approx.	kg	27
Noise level * ca. without housing with housing	 dB(A) dB(A)	 65 61

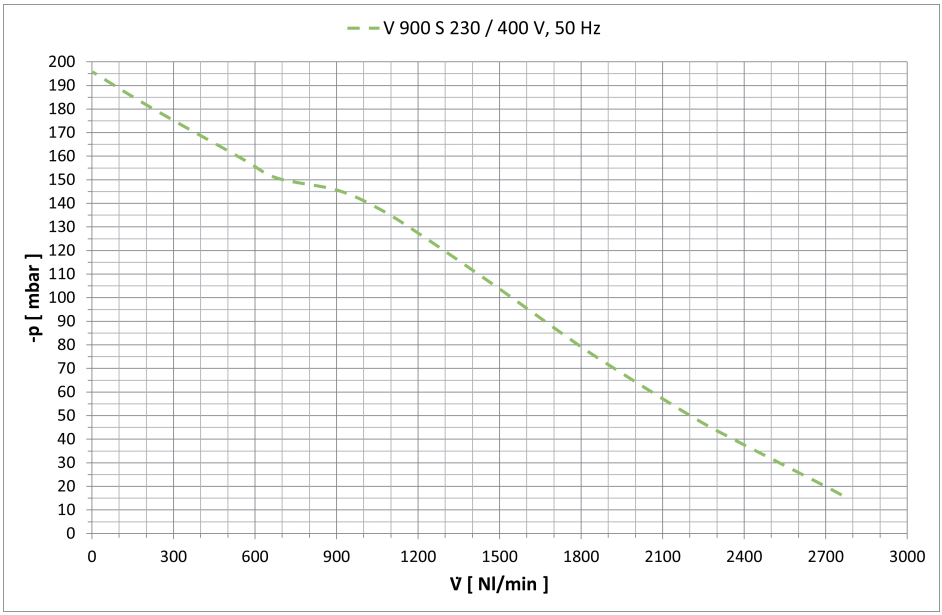
* Noise level in accordance with ISO 3746

Ambient conditions during storage and transport		
Temperature	°C	-10 - +60
Relative humidity	%	< 95

Ambient conditions during operation		
Temperature	°C	+10 - +40
Relative humidity	%	< 70

Classification	
Medical Device Class (MDR)	IIa

Characteristic curve in accordance with EN ISO 10637



Characteristic curve for: 7131-02

4.6 V 1200 S

Electrical data		7136-02	
Rated voltage	V	230, 3~	400, 3~
Mains frequency	Hz	50	
Active power	kW	1.8	1.8
Nominal current	A	≤ 6.2	≤ 3.6
Start-up current	A	~ 43	~ 25
Motor protection *	A	6.3	4.0
Type of protection		IP 24	
Protection class		I	

* Max. permitted setting

Connections		
Suction connection (outside)	mm	∅ 50
Exhaust air connection (external)	mm	∅ 50

Classification in accordance with EN ISO 10637		
Classification based on separation of solids and liquids		Dry suction system
Classification based on flow rate		Type 1

General data		
Max. number of users		4
Duty cycle	%	100 (S1)
Heat generation rate	MJ/h	6.5
Dimensions (H x W x D)	cm	52 x 41 x 43
Weight, approx.	kg	27
Noise level * ca. without housing with housing	 dB(A) dB(A)	 65 61

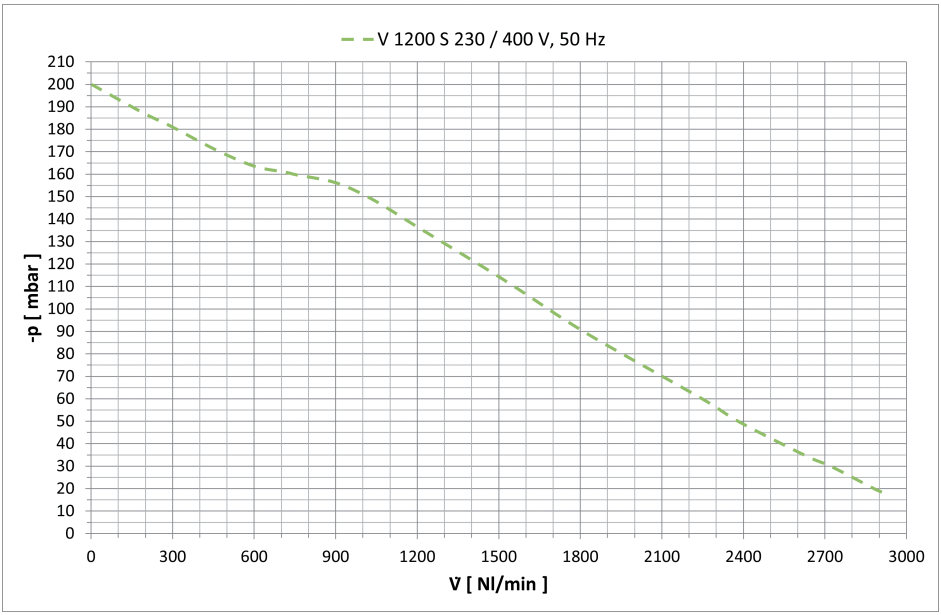
* Noise level in accordance with ISO 3746

Ambient conditions during storage and transport		
Temperature	°C	-10 - +60
Relative humidity	%	< 95

Ambient conditions during operation		
Temperature	°C	+10 - +40
Relative humidity	%	< 70

Classification	
Medical Device Class (MDR)	IIa

Characteristic curve in accordance with EN ISO 10637

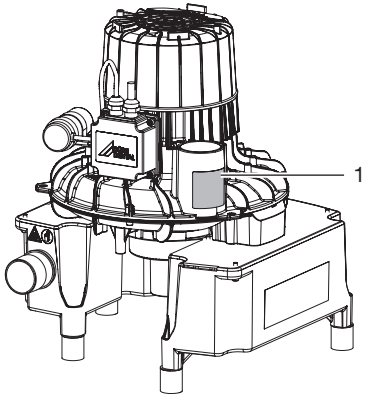


Characteristic curve for: 7136-02

4.7 Type plate

V 600

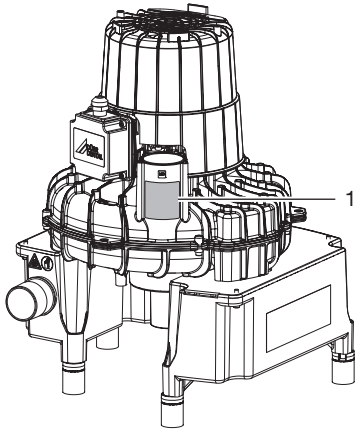
The type plate is located on the upper part of the turbine housing.



1 Type plate

V 900 S, V 1200 S

The type plate is located on the upper part of the turbine housing.

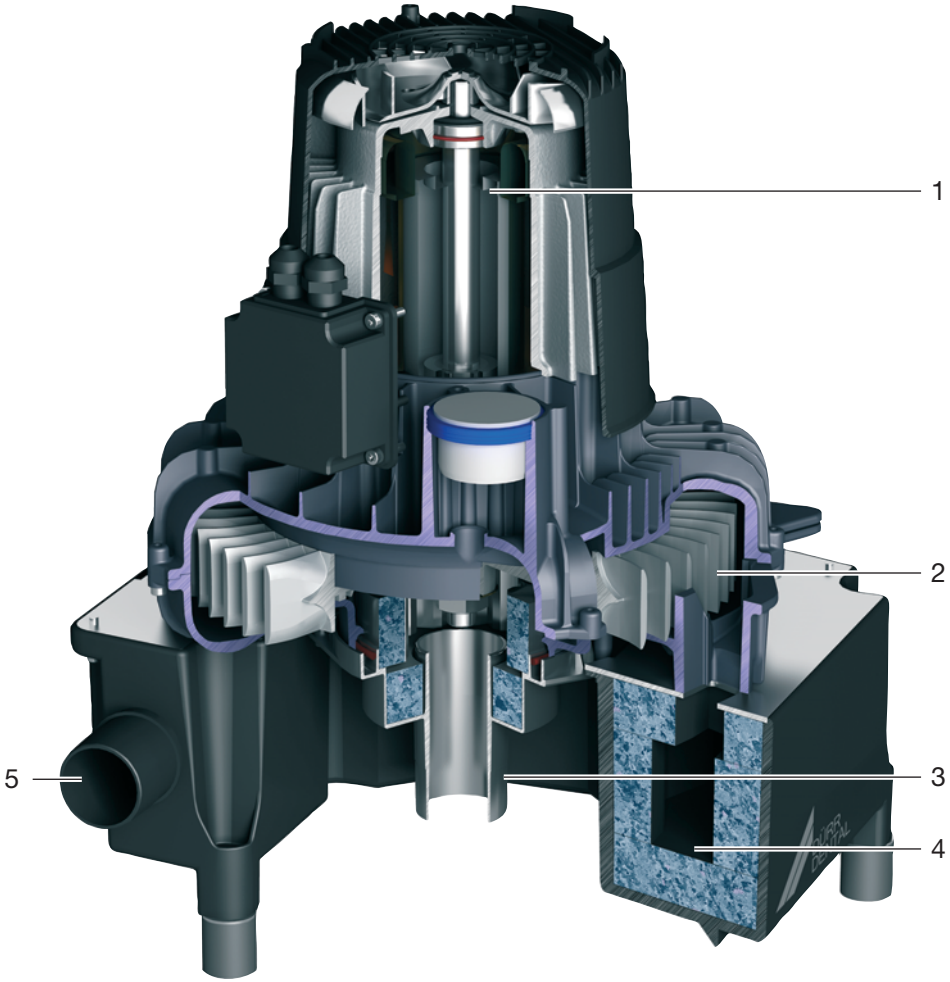


1 Type plate

4.8 Evaluation of conformity

This device has been subjected to conformity acceptance testing in accordance with the current relevant European Union guidelines. This equipment conforms to all relevant requirements.

5 Function



1 Motor
2 Turbine wheel
3 Intake nozzle
4 Exhaust air muffler
5 Exhaust air connection

The V-suction unit is suitable for use in dry air suction systems. The advantage of this system is that the suction unit, regardless of the actual connection layout, can be installed in any available and suitable room (including upper floors or basements). The necessary air flow and vacuum are generated by a rapidly rotating impeller.
When an appropriate vacuum for the machine is applied, approx. 300 l/min of air is sucked in through the suction cannula.

On the vacuum side the V-suction unit is equipped with a condensation separator that collects any condensation arising within the pipe system and transports it away to the outside. An auxiliary air valve in the condensation separator protects the suction unit against overheating and provides uniform suction power.

The auxiliary air valve is matched to the suction unit. It must not be modified or replaced with an auxiliary air valve with a different setting.

The exhaust air from the suction unit should be guided out of the building (via the roof where possible). We recommended the installation of a bacteria filter in the exhaust air line. In addition, it is possible to install a noise-reducing muffler in the exhaust air line in order to reduce the amount of noise generated by the unit and by the air flow.

Assembly

6 Requirements

The unit can be installed either on the same level as the surgery or on a floor below.

Further information can be found in our suction planning information leaflet.
Order number9000-617-03/..

6.1 Installation/setup room

The room chosen for set up must fulfil the following requirements:
- Closed, dry, well-ventilated room
- Should not be a room made for another purpose (e. g. boiler room or wet cell)
- When installing in a cabinet the inlet and outlet ventilation slots must be present; minimum free cross-section at least 120 cm^2.
- Forced ventilation (fan) must be provided if there is a risk that the recommended room air temperature could be exceeded. The air flow performance must be at least 2 m^3/min.
- Do not cover cooling slots or openings with housing installations; ensure sufficient clearance to the openings to permit sufficient cooling.

6.2 Setup options

The following options for setting up the unit are available:
- Wall installation using a Dürr Dental wall mounting
- In a ventilated cabinet
- In a Dürr Dental noise reducing housing

6.3 Pipe materials

Only use vacuum-sealed HT-waste pipes manufactured from the following materials:
- Polypropylene (PP),
- Chlorinated polyvinyl chloride (PVC-C),
- Plasticizer-free polyvinyl chloride (PVC-U),
- Polyethylene (PE).

The following materials must not be used:
- Acrylonitrile-butadiene-styrene (ABS),
- Styrene copolymer blends (e.g. SAN + PVC).

6.4 Hose materials

For waste connections and suction lines only use the following hose types:
- Flexible spiral hoses made of PVC with integrated spiral or equivalent hoses
- Hoses that are resistant to dental disinfectants and chemicals

Plastic hoses will display signs of ageing over time. Therefore, they should be inspected regularly and replaced as necessary.

The following types of hoses must not be used:
- Rubber hoses
- Hoses made completely of PVC
- Hoses that are not sufficiently flexible

6.5 Information about electrical connections

- Ensure that the electrical connections to the mains power supply are established in accordance with current valid national and local regulations and standards governing the installation of low voltage units in medical facilities.
- Install an all-pole disconnect switch with a contact opening width of at least 3 mm in the electrical connection to the mains power supply.
- Observe the current consumption of the devices that are to be connected.

Electrical fusing

LS switch 16 A, characteristic B, C and D in accordance with 60898.

6.6 Information about connecting cables

The diameter of the connections depends on the current consumption, length of line and the ambient temperature of the unit. Information concerning the current consumption can be found in the Technical Data supplied with the particular unit to be connected.
The following table lists the minimum diameters of the connections in relation to the current consumption:

Current consumption of unit [A]	Cross-section [mm^2]
> 10 and < 16	1.5
> 16 and < 25	2.5

Current consumption of unit [A]	Cross-section [mm²]
> 25 and < 32	4
> 32 and < 40	6
> 40 and < 50	10
> 50 and < 63	16

Mains supply cable

Installation type	Line layout (minimum requirements)
Fixed installation	– Plastic sheathed cable (e.g. type NYM-J)
Flexible	– PVC flexible line (e.g. H05 VV-F) or – Rubber connection (e.g. H05 RN-F or H05 RR-F)

Control cable

24 V protective low voltage for:
– Hose manifold
– Place selection valve
– Spittoon valve

Installation type	Line layout (minimum requirements)
Fixed installation	– Shielded sheathed cable (e.g. (N)YM (St)-J)
Flexible	– PVC data cable with shielded cable sheathing, as used for telecommunications and IT processing systems (e.g. type LiYCY) or – Lightweight PVC control cable with shielded cable sheathing

Connect the shielding of the cables in accordance with the regulations.

7 System components

The system components listed below are required or recommended for various procedures or for installation.

7.1 Control box

The unit is connected via a control box. The control box is either included in the scope of delivery or must be ordered separately. In some units, the control system is built in.

7.2 Exhaust air filter

For hygienic reasons, we recommend the installation of a bacteria filter in the exhaust air line. If the unit is installed in the surgery and the exhaust air cannot be discharged to the outdoors, it is essential to install a bacteria filter. Depending on the type and condition of the bacteria filter, it will need to be replaced every 1-2 years at the latest.

The separation integrated in the system does not retain bacteria; this is why we recommend installing a suitable filter in the exhaust air system.

7.3 Noise reduction

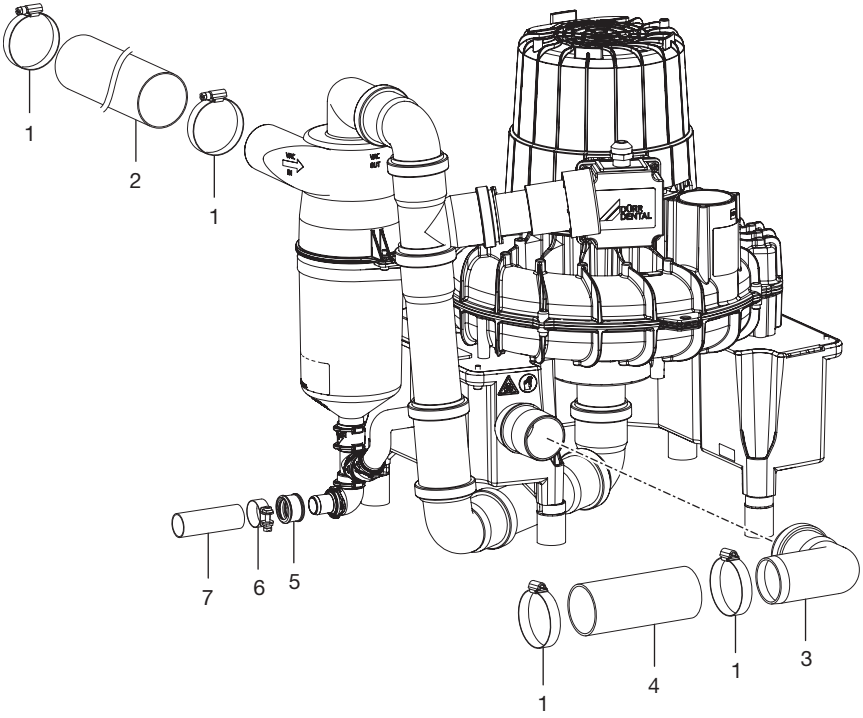
If the noise level from the exhaust air vent or the flow noise generated is too high, noise reduction can be installed in the exhaust air line.

8 Installation

The actual connection can vary depending on the chosen installation option. The connection shown is only an example.

8.1 Installation and routeing of hoses and pipes

– Establish connections between the pipe system and the unit using the flexible hoses supplied. This will prevent vibrations from being transmitted to the pipe system.
– The connection between the pipe line and unit suction connection should be kept as short as possible and straight, without bends.
– Install the drain hoses with a downward gradient so that the waste water can drain off.
– Waste water connections must be implemented in accordance with applicable local and national regulations.



1 Hose clip 40-60 mm
2 Suction hose ∅ 50 mm
3 Elbow DN 50 / 87°
4 Waste air pipe (aluminium)∅ 50 mm inside
5 Hose sleeve
6 Hose clamp ∅ 28 mm
7 Waste water hose ∅ 20 mm (internal)

8.2 Electrical connections

WARNING

Electric shock due to incorrectly connected device

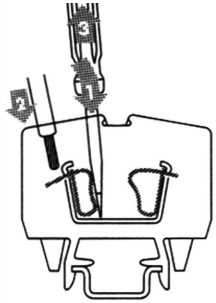
› Never install a mains plug instead of the fixed connection.

NOTICE

Short circuit due to defective lead

› Do not route wires near hot surfaces.

– Before connecting, check to make sure that the mains voltage matches the voltage specifications on the type plate.
– Only connect the unit to an original control box.
– Connect the power supply line from the control box to the appropriate terminals in the motor terminal box.



1. Insert screwdriver into terminal strip to open spring clamp.
2. Insert stranded wire into spring clamp.
3. Remove screwdriver.

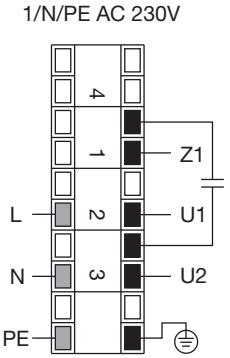


Fig. 1: Terminal assignment for 1-phase units

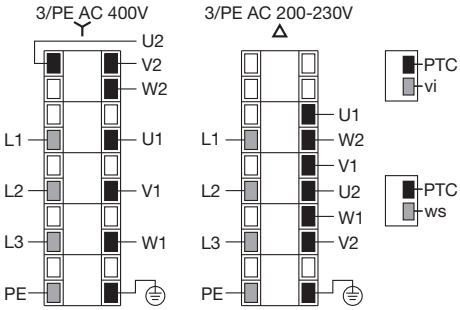


Fig. 2: Terminal assignment for 3-phase units

8.3 Testing suction power after installation

The following procedure is recommended for measuring the maximum negative pressure of the suction system for comparison with the performance curve:

1. Lift a suction hose off a treatment unit and connect a pressure gauge (relative pressure). The connection adapter to the pressure gauge must be leak tight.
2. Read the displayed negative pressure and compare it against the performance curve.

9 Commissioning

In many countries technical medical products and electrical devices are subject to regular checks at set intervals. The owner must be instructed accordingly.

1. Turn on the unit power switch or the main surgery switch.
2. Carry out a function check of the system.
3. Check all connections for leak tightness.
4. Carry out an electrical safety check in accordance with applicable regulations (e.g. regulations concerning set up, operation and application of medical devices) and record the results as appropriate, e.g. in the technical log book.
5. Carry out and document the instruction and handover for the unit.

A sample handover report is included in the attachment.

Usage

10 Disinfection and cleaning

NOTICE

Device malfunctions or damage due to use of incorrect media

Guarantee claims may become invalid as a result.

- Do not use any foaming agents such as household cleaning agents or instrument disinfectants.
- Do not use abrasive cleaners.
- Do not use agents containing chlorine.
- Do not use any solvents like acetone.

As a general rule use:
– for disinfection and cleaning:
Orotol plus or Orotol ultra
– for cleaning:
MD 555 cleaner

Only these products have been tested by Dürr Dental.
When using prophy powders, water-soluble Lunos Prophy Powders are recommended in order to protect the suction systems (Dürr Dental).

10.1 After every treatment

1. Aspirate a glass of cold water through the large and the small suction hoses. Do this even if only the small suction hose was actually used during treatment.

Suction through the large suction hose causes a large amount of air to be drawn up, thereby considerably increasing the cleaning effect.

10.2 Daily after the end of treatment

After higher workloads before the midday break and in the evening

The following are required for disinfection/cleaning:

✓ Non-foaming disinfectant/cleaning agent that is compatible with the materials.
✓ Unit care system, e.g. OroCup

1. To pre-clean, suck up 2 litres of water with the care system.
2. Aspirate the disinfection/cleaning agent with the care system.

10.3 Once or twice a week before the midday break

Under harsher conditions (e.g. hard water or frequent use of prophy powders) 1x daily before the midday break

The following are required for cleaning:

✓ Special non-foaming detergent for suction units that is compatible with the materials.
✓ Unit care system, e.g. OroCup

1. To pre-clean, suck up 2 litres of water with the care system.
2. Aspirate the cleaning agent with the care system.
3. Rinse with ca. 2 l water after the application time.

11 Maintenance

All maintenance work must be performed by a qualified expert or by one of our Service Technicians.

WARNING

Infection due to contaminated unit

- Clean and disinfect the suction before working on the unit.
- Wear protective equipment when working (e. g. impermeable gloves, protective goggles and mouth and nose protection).

Prior to working on the unit or in case of danger, disconnect it from the mains.

Maintenance interval	Maintenance work
Every 1-2 years	› Replace the exhaust air filter (where fitted). *
Every 2 years	› Check the waste valve on the condensation separator for correct operation and replace it if necessary. * › Check the auxiliary air valve for correct operation and clean/replace it as required. *

* Only to be performed by service technicians.

Troubleshooting

12 Tips for operators and service technicians

Any repairs exceeding routine maintenance may only be carried out by qualified personnel or our service.

WARNING

Infection due to contaminated unit

- Clean and disinfect the suction before working on the unit.
- Wear protective equipment when working (e. g. impermeable gloves, protective goggles and mouth and nose protection).

Prior to working on the unit or in case of danger, disconnect it from the mains.

Error	Possible cause	Remedy
Device does not start	No mains voltage	› Check the mains supply voltage. * › Check the fuses and replace if necessary. *
	Undervoltage	› Measure the supply voltage; call an electrician if necessary. *
	Motor protection switch set too low	› Measure current, set the motor protection switch to the measured value plus safety margin. *
	Motor protection switch defective	› Check the motor protection switch; replace if defective. *
	Capacitor defective	› Measure capacitance and replace if necessary. *
	Turbine is blocked by solid particles or sticky soiling	› Disassemble the unit and clean the turbine. *
The unit generates unusual noises	Solid particles in the turbine chamber	› Disassemble the unit and clean the turbine and housing. *
Water leaking from the exhaust air connection	Foam in turbine due to use of incorrect cleaning and disinfectant agents	› Use non-foaming cleaning and disinfectant agents.
	Build-up of condensation in the exhaust air line	› Check the pipe system; avoid over-cooling. *

Error	Possible cause	Remedy
Suction performance too low	Coarse filters in system clogged (e.g. at separator devices)	› Clean coarse filters.
	Leak in the suction line	› Check and if necessary establish leak-tightness of suction system and connections. *
	Mechanical sluggishness of turbine caused by soiling	› Disassemble the unit and clean the turbine. *

* Only to be done by service technicians.

13 Transporting the unit

WARNING

Infection due to contaminated unit

- Disinfect the unit before transport.
- Close all media connections.

Wear protective equipment to avoid any risk of infection (e.g. liquid-tight protective gloves, protective goggles, face mask).

1. Prior to disassembly, clean and disinfect the suction unit and the unit via aspiration of a suitable disinfectant approved by the manufacturer.
2. Disinfect a defective unit using a suitable surface disinfection agent.
3. Seal all connections with sealing caps.
4. Pack the unit securely in preparation for transport.

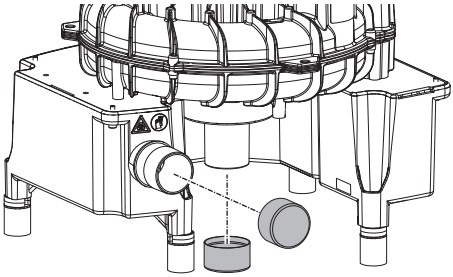


Fig. 3: V 600, V 900 S, V 1200 S

1 Sealing cap

Appendix

14 Handover record

This document confirms that a qualified handover of the medical device has taken place and that appropriate instructions have been provided for it. This must be carried out by a qualified adviser for the medical device, who will instruct you in the proper handling and operation of the medical device.

Product name	Order number (REF)	Serial number (SN)

- ☐ Visual inspection of the packaging for any damage
- ☐ Unpacking the medical device and checking for damage
- ☐ Confirmation of the completeness of the delivery
- ☐ Instruction in the proper handling and operation of the medical device based on the operating instructions

Notes:

Name of person receiving instruction: **Signature:**

Name and address of the qualified adviser for the medical device:

Date of handover: **Signature of the qualified adviser for the medical device:**

15 Country representatives

Country		Address
GB	UK CA XXXX	**UK Responsible Person:** Duerr Dental (Products) UK Ltd. 14 Linnell Way Telford Way Industrial Estate Kettering, Northants NN 16 8PS
UA	UA.TR.099	**Уповноважений представник в Україні:** Приватне підприємство “Галіт” вул. 15 квітня, 6Є, с. Байківці, Тернопільський р-н, 47711, Україна тел.: 0800 502 998; +38 050 338 10 64 www.galit.te.ua; e-mail: office@galit.te.ua Виробник: Дюрр Дентал ЕсЕ Хьопфігхаймер Штрассе 17, Д-74321 Бітігхайм-Біссінген, Німеччина email: info@duerrdental.com
CN		**备**案人/生**产**企**业**： DÜRR DENTAL SE 德国迪珥**齿**科股份公司 住所/生**产**地址： Höpfigheimer Str. 17, 74321 Bietigheim-Bissingen, Germany **联**系方式： **电话**： + 49 7142 705-0 **邮**箱： info@duerrdental.com 网址： www.duerrdental.com 代理人/售后服**务单**位：迪珥医**疗**器械（上海）有限公司 住所：上海市**长**宁区天山路 641 号 2 号楼 (20 幢) 303 室 **联**系方式： **电话**：+ 86 21 6381 0270 **传**真：+ 86 21 6381 0290 **邮**箱：info@duerr.cn 网址：http://www.duerrdental.com

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