

EN Instruction Manual

ES Manual de usuario

DE Bedienungsanleitung

FR Mode d'emploi

RU Руководство по эксплуатации

PL Instrukcja obsługi

AR كتيب التعليمات

FA دستورالعمل راهنما

www.rossmax.com



Warranty Registration (Must be completed within 10 days of purchase)

Customer Name: _____ **City:** _____
Street Address: _____ **ZIP code:** _____
State: _____ **ZIP code:** _____
Telephone: _____ **E-mail address:** _____

Gender: Male Female **Age:** _____

Product Information

Date of purchase: _____

Store where purchased: _____

Price Paid (excl. Tax): _____

This instrument is covered by a 3 year guarantee from the purchase date. The guarantee is valid only on presentation of the guarantee card completed by the dealer conforming purchase date or the receipt. Nebulizer Components are not included. Opening or altering the instrument invalidates the guarantee. The guarantee does not cover damage, accident or non-compliance with the instruction manual. Please contact Rossmax Service or distributors.

ISO 9001

ISO 14001

ISO 13485

ISO 27001

ISO 22700

ISO 22703

ISO 22704

ISO 22705

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Introduction
Thank you for purchasing a NE100 Compressor Nebulizer. With proper care and use, your nebulizer will provide you with many years of reliable treatments. This unit operates on standard AC power. Treatments are delivered quickly, safely and conveniently, making this unit ideal for all ages. We encourage you to thoroughly read this guide-book to learn about the features of your nebulizer. Your compressor nebulizer should be used under the supervision of a licensed physician and/or a respiratory therapist. Together with your physician and/or pharmacist, you can feel comfortable and confident knowing that you are obtaining the most effective inhalation treatments for your respiratory condition.

NOTE: Your nebulizer is intended for use in treatment of asthma, COPD and other respiratory ailments in which an aerosolized medication is required during therapy. Please consult with your physician and/or pharmacist to determine if your prescription medication is approved for use with this nebulizer. For type, dose, and regime of medication follow the instructions of your doctor or respiratory therapist.

This device fulfills the provision of the EC directive 93/42/EEC (Medical Device Directive) and the European Standard EN 13544-1:2007+A1:2009 (Respiratory Therapy Device Unit - Part 1: Nebulizing systems and their components).

PLEASE read this manual carefully before use and be sure to keep this manual.

Cautions
Please use general safety precautions when operating your nebulizer. This unit should be used only for its intended purpose as described in this guidebook and with medications only under the supervision and instruction of your physician. Do not use the device in anesthetic or ventilator breathing circuits.

Product cautions

READ THE FOLLOWING BEFORE USING

- Do not use this unit while bathing or showering.
- Do not immerse the unit in liquid.
- Do not handle the unit of power cord with wet hands.
- Do not use the unit in liquid.
- Do not use while bathing.
- Do not reach for anything that has fallen into water immediately upon the unit.
- Do not use the unit if it has any damaged parts (including plug). If it has been submerged in water or dropped, promptly send the unit for examination and repair.
- The unit should not be used where flammable gas, oxygen or aerosol spray products are being used.
- Keep the air vents open. Do not place the unit on a soft surface where the openings can be blocked.
- If the medication cup is empty, do not attempt to operate the unit.
- If any abnormality occurs, discontinue to use until the unit has been examined and repaired.
- The unit should not be left unattended while plugged in.
- Do not tilt or shake the unit when in operation.
- Disconnect the unit from the electrical outlet before cleaning, filling and after each use.
- Do not use attachments unless recommended by the manufacturer.
- Do not disassemble or attempt to repair the unit.
- Do not use the device in anesthetic or ventilator breathing circuits.

Operating cautions

- Close adult supervision is highly recommended when the unit is used by children or invalids.
- Keep your eyes away from the output of medication mist.
- The maximum capacity of the medication cup is 5 ml and should not be overfilled.
- Do not use this unit while operating a vehicle.
- If any discomfort or abnormality occurs, stop using the unit immediately.
- Do not use the device if the air tube is bent.
- Pentamidine is not an approved medication for use with this device.

Storage cautions

- Do not store the unit in direct sunlight, high temperature or humidity.
- Keep the unit out of reach of small children.
- Always keep the unit unplugged while not in use

Cleaning cautions

- Check air filter, nebulizer, mouthpiece and any other optional component before each use. Dirty or worn parts should be replaced.
- Do not immerse the unit in water. It may damage the unit.
- Disconnect the unit from the electrical outlet before cleaning.
- Clean all necessary parts after each use as instructed in this guidebook.
- Always dispose of any remaining medication in the medication cup after each use. Use fresh medication in the medication cup.
- Do not store the air tube with moisture or medication remaining in the air tube. This could result in infection as a result of bacteria.

MEDICAL DISCLAIMER:

This manual and product are not meant to be a substitute for advice provided by your doctor or other medical professional. Don't use the information contained herein or this product for diagnosing or treating a health problem or prescribing any medication. If you have or suspect that you have a medical problem, promptly consult your doctor.

Durable periods are as follows, provided the product is used to nebulize 2ml of medication 2 times a day for 8 minutes each time at room temperature (25°C). Durable period may vary depending on usage environment.

Main unit	5 years
Nebulizer Kit	1 year
Air Tube, Mouthpiece	1 year
Air Filter	60 days
Adult and child masks	5 years

Product specifications

Power	AC 230V/50Hz or AC 220V/60Hz or AC 110V/60Hz
Power Consumption	≤ 130W
Sound Level	≤ 60 dBA (1 meter away from NE100)
Compressor Pressure Range	≥ 29 psi (200 kPa)
Operating Pressure Range	≥ 15 psi (103 kPa)
Operating Flow Range	≥ 3.5 lpm
Operating Temperature Range	10°C to 40°C (50°F to 104°F)
Operating Humidity Range	10 – 90% RH
Operating Atmospheric Pressure Range	700-1060 hPa
Storage Temperature Range	-20°C to 60°C (-4°F to 140°F)
Storage Humidity Range	10 – 90% RH
Dimension (L x W x H)	204.5mmx148mmx108mm (8.05x5.83x4.25 inches)
Weight	1360g (without accessories)
Medication Capacity	5ml(cc)
Particle Size (MMAD)	≤ 3.0µm
Average Nebulization Rate	≥ 0.3 ml/min
Standard Accessories	Nebulizer Kit, Air Tube, Mouthpiece, Filters (5pcs), Adult and child masks

^aSubject to technical modification without prior notice.
^bPerformance may vary with drugs such as suspensions or high viscosity. See drug supplier's data sheet for further details.

A. Product identification			
1. Nebulizer Kit	2. Nozzle	3. Angled Mouthpiece	
4. Air Filter	5. Child Mask	6. Adult Mask	
7. Air Tube	8. Air Tube	9. Air Filter Slot	
10. Air Intake	11. Power Cord	12. Power Switch	

B. Assembling your nebulizer kit
Follow the cleaning instructions in this guidebook under "Cleaning technique" prior to using your nebulizer for the first time or after it has been stored for an extended period of time.

REMEMBER: Always unplug the compressor and make sure the power-switch is turned to the "OFF" position before cleaning, assembling and before or after each use.

- Place the compressor on a flat, stable surface within reach.
- Gently twist and pull straight up on the lid of the nebulizer to separate into two parts (medication cup and cover).
- The stem inside the medication cup inserts into the tube of the nozzle.
- Add the prescribed amount of medication in the medication cup.
- Reassemble the nebulizer by carefully twisting the medication cup and cover together. Make sure that the two parts fit securely.

WARNING: The symbol on this product means that it's an electronic product and following the European directive 2012/19/EU the electronic products have to be disposed on your local recycling centre for safe treatment.

C. Operating your nebulizer

The nebulizer is operate at up to a 45° angle. If the angle is greater than 45°, no aerosol will be generated.

- Attach one end of the air tube connector to the air output.
- Carefully attach the opposite end of the air tubing connector to the stem at the base of the nebulizer kit.
- Attach the angled mouthpiece or mask to the top of the nebulizer.
- The capacity of the medication cup is 2-5 ml.

NOTE: A 30-minute interval is recommended after each use. The compressor will automatically shut off if it becomes overheated. If it when this happens, immediately:

- Press the power-switch to the "OFF" position.
 - Unplug the power cord from the outlet.
 - Allow the motor to cool for 30 minutes before using the nebulizer again.
- Before restarting the unit, make sure that the air vents are not obstructed.

Operating your nebulizer

After every use:

- Unplug the unit from the power source.
- Allow the unit to completely cool.
- Carefully detach the air tubing from the nebulizer and pour out any remaining medication.

4. Follow the cleaning procedures provided in this guidebook.

D. Cleaning procedures

Rinsing technique (performed after each treatment or before first use):

- Make sure that the power-switch has been turned to the "OFF" position and the unit has been disconnected from the power source.
- Disconnect the air tube from the nebulizer device.
- Gently twist and pull up the cover of the nebulizer kit to open and separate.
- Rinse the nebulizer kit and components with hot tap water.
- Dry with clean towels or completely air dry.
- Reassemble the nebulizer kit.

NOTE: For the first time cleaning or after the unit has been stored for an extended period of time, thoroughly clean all components, except the air tube. The nebulizer kit is dishwasher safe.

Cleaning the compressor

Wipe the compressor daily using a soft cloth.

NOTE: Any other form of cleaning or cleaning agents may damage the finish of the unit.

E. Changing the air filter

It is important to change the air filter approximately when the air filter turns gray. It is recommended to change air filter every 2 months.

- Remove the air filter cover by gently pulling forward.
 - Discard the gray filter.
 - Replace with a new, clean air filter.
 - Securely re-attach the air filter cover to the unit.
- NOTE:** Air filters cannot be cleaned or washed. Only NE100 air filters can be used. Do not substitute alternate material such as cotton. Do not operate without an air filter.

Troubleshooting

- If any abnormality occurs during use, please check and correct the following:
- Unit does not operate when power switch is pressed. Check the AC connection to the outlet.
 - No misting or low rate of misting:
 - Check that there is medication in the nebulizer cup.
 - Check the main unit if there is any physical damage.
 - Check the position of the nozzle inside the nebulizer.
 - Make sure that air tube and other components are properly attached.
 - Check the air filter and replace if necessary.

Protection against electric shock:

- Class II equipment

Type BF applied parts:

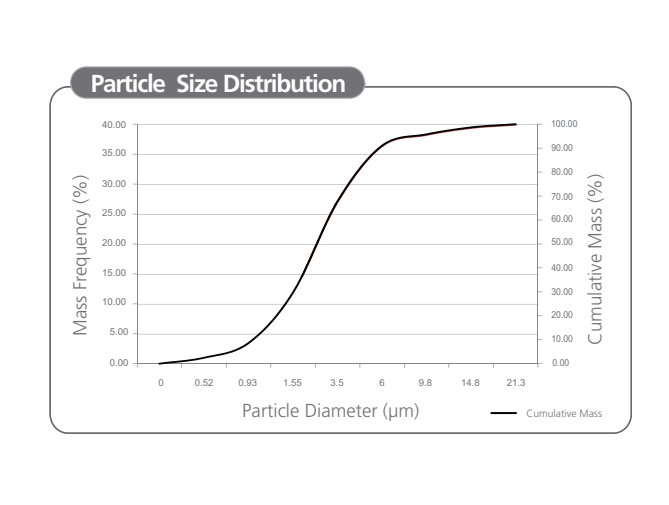
- III and II parts and accessories

Protection against harmful ingress of water and particulate matter:

- IP21

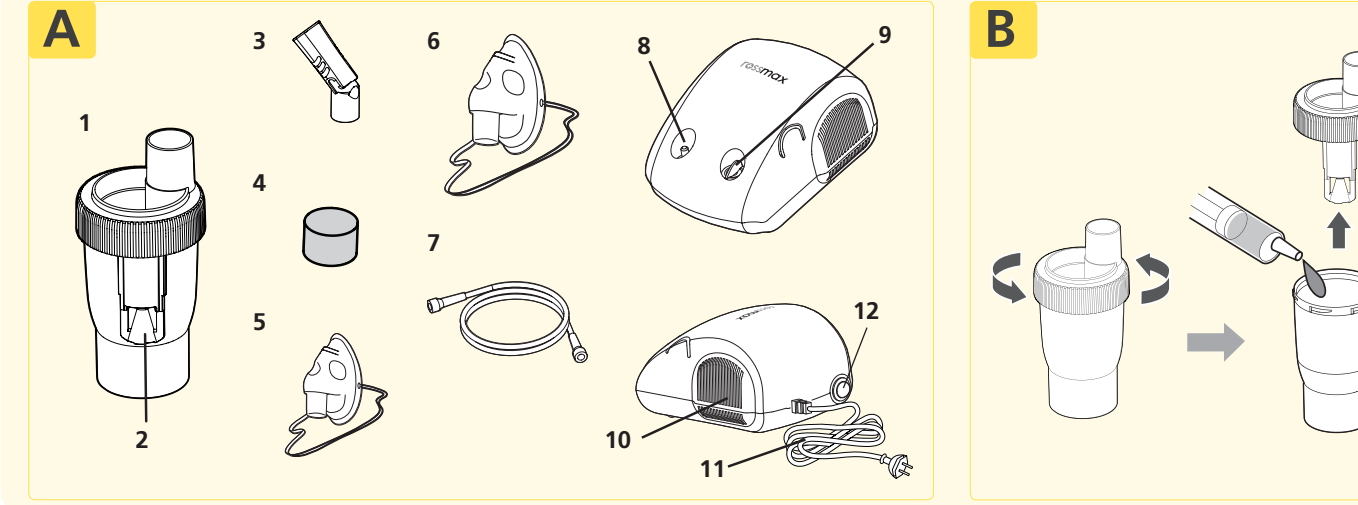
Degree of safety in the presence of flammable anesthetics or oxygen:

- No APAPG (not suitable for use in the presence of flammable anesthetics or oxygen).



Guidance and manufacturer's declaration-electromagnetic emissions			
The NE100 is intended for use in the electromagnetic environment specified below. The customer or the user of the NE100 should assure that it is used in such an environment.			
Immunity test	EC 60801 test level	Completion level	Electromagnetic environment-guidance
RF emissions CISPR 11	Group 1	Class B	The NE100 uses RF energy only for its internal function. Therefore, as RF emissions are very low and are not likely to cause any interference in nearby electronic equipment.
RF emissions CISPR 11	Class B	Class A	The NE100 is suitable for use in all establishments. Harmonic emissions IEC 61000-3-2
RF emissions CISPR 11	Class B	Class A	The NE100 is suitable for use in all establishments. Harmonic emissions IEC 61000-3-2
Voltage fluctuations/flicker emissions IEC 61000-3-3	Compliance		those directly connected to the public low-voltage power supply network that supplies buildings used for domestic purposes.
Guidance and manufacturer's declaration-electromagnetic immunity			
The NE100 is intended for use in the electromagnetic environment specified below. The customer or the user of the NE100 should assure that it is used in such an environment.			
Immunity test	EC 60801 test level	Completion level	Electromagnetic environment-guidance
Electrostatic discharge IEC61000-4-2	±6 kV contact ±8 kV air	±6 kV contact ±8 kV air	Floors should be wood, concrete or ceramic tile. If floors are covered with synthetic material, the relative humidity should be at least 30%.
Electrical fast transient/burst supply lines IEC 61000-4-4	±2kV for power supply lines ±2kV for input/output lines	±2kV for power supply lines Not applicable	Mains power quality should be that of a typical commercial or hospital environment.
Surge IEC 61000-4-5	±1kV differential mode ±2kV line-to-earth	±1kV differential mode Not applicable	Mains power quality should be that of a typical commercial or hospital environment.
Voltage dips IEC 61000-4-11	<5% UTI>95% dip in short interruption 40% <UTI>80% dip in long variations on power supply	<5% UTI>95% dip in short interruption 40% <UTI>80% dip in long variations on power supply	Mains power quality should be that of a typical commercial or hospital environment. The use of the NE100 requires continued operation during power mains interruptions. It is recommended that the NE100 be powered by an uninterruptible power supply (UPS) or a battery.
Power IEC 61000-4-7	3AIm	3AIm	The NE100 power frequency magnetic fields should be at levels characteristic of a typical location in a typical commercial or hospital environment.
NOTE: UTI is the a.c. mains voltage prior to application of the test level.			
Guidance and manufacturer's declaration-electromagnetic immunity			
The NE100 is intended for use in the electromagnetic environment specified below. The customer or the user of the NE100 should assure that it is used in such an environment.			
Immunity test	EC 60801 test level	Completion level	Electromagnetic environment-guidance
Portable and mobile RF communications equipment should be used no closer to any part of the NE100 including cables than the recommended separation distance calculated from the equation applicable to the frequency of the transmitter. Recommended separation distance:			
Where P is the maximum output power rating of the transmitter in (W) according to the transmitter manufacturer and d is the distance in metres (m).			
Field strengths from fixed RF transmitters, as determined by an electromagnetic site survey, ^a should be less than the compliance level in each frequency range. ^b Interference may occur if the NE100 is not in normal operation. If abnormal performance is observed, additional measures may be necessary, such as re-orienting or relocating the NE100.			
b. Over the frequency range 150 kHz to 80 MHz, field strengths should be less than 3 µV/m.			
NOTE1: At 80 MHz and 800 MHz, the higher frequency range applies.			
NOTE2: These guidelines may not apply in all situations. Electromagnetic propagation is affected by absorption and reflection from structures, objects and people.			
^a Field strengths from fixed transmitters, such as base stations for radio (cellular/cordless) tele-phones and land mobile radios, amateur radio, AM and FM radio broadcast and TV broadcast cannot be predicted theoretically with accuracy. To assess the electromagnetic environment due to fixed RF transmitters, an electromagnetic site survey should be considered. If the measured field strength in the location in which the NE100 is used exceeds the applicable RF compliance level above, the NE100 should be observed to verify normal operation. If abnormal performance is observed, additional measures may be necessary, such as re-orienting or relocating the NE100.			
^b Over the frequency range 150 kHz to 80 MHz, field strengths should be less than 3 µV/m.			

Recommended separation distance between portable and mobile RF communications equipment and the NE100			
Conducted RF IEC 61000-1-4	3 Vrms 150 kHz to 80 MHz	3 Vrms 150 kHz to 80 MHz	3 Vrms 150 kHz to 80 MHz
Radiated RF IEC 61000-1-3	3 V/m 80 MHz to 2.5 GHz	3 V/m 80 MHz to 2.5 GHz	3 V/m 80 MHz to 2.5 GHz
Recommended separation distance according to frequency of transmitter (m)			
Power (W)	d=1.2√P	d=1.2√P	d=1.2√P
0.01	0.12	0.12	0.23
0.1	0.38	0.38	0.73
1	1.2	1.2	2.3
10	3.8	3.8	7.3
100	12	12	23
For transmitters rated at a maximum output power not listed above, the recommended separation distance in metres (m) can be estimated using the equation applicable to the frequency of the transmitter. Where P is the maximum output power rating of the transmitter in watts (W) according to the transmitter manufacturer.			
NOTE1: At 80 MHz and 800 MHz, the separation distance for the higher frequency range applies.			
NOTE2: These guidelines may not apply in all situations. Electromagnetic propagation is affected by absorption and reflection from structures, objects and people.			



Es Español

Introducción
Gracias por comprar el nebulizador de compresor NE100. Con el cuidado y uso adecuado, su nebulizador le proporcionará tratamientos fiables por muchos años. La unidad funciona con corriente alterna estándar. Los tratamientos se conducen de modo rápido, seguro y conveniente, haciendo que la unidad sea ideal para cualquier edad. Recomendamos que lea este manual de usuario atentamente para llegar a conocer las características de su nebulizador. Su nebulizador de compresor debería utilizarse bajo la supervisión de un médico y/o terapeuta respiratorio licenciado. Junto con su médico o su terapeuta respiratorio, podrá sentirse confortable y confiante sabiendo que está obteniendo los tratamientos de inhalación más efectivos para su enfermedad respiratoria.

NOTA: Su nebulizador se ha concebido para su uso en el tratamiento de asma, EPOC u otras enfermedades respiratorias, en las cuales se requiere una medicación aerosolizada durante la terapia. Por favor consulte a su médico y/o farmacéutico para determinar si su medicación recetada está autorizada para el uso con este nebulizador. Con respecto al tipo, a la dosis y al régimen de la medicación, observe las instrucciones de su médico o terapeuta respiratorio.

Este aparato satisface las exigencias de la Directiva 93/42/CEE (Directiva de Productos Sanitarios) de la CE y la Norma Europea EN 13544-1:2007+A1:2009 Equipos de terapia respiratoria - Parte 1: Sistemas de nebulización y sus componentes.

IP21
Por favor, lea este manual cuidadosamente antes del uso. Por favor, fíjese en el cuadro de advertencias.

Precauciones

Por favor, tome las precauciones de seguridad generales al estar operando su nebulizador. Esta unidad sólo deberá usarse para el fin descrito en este manual de usuario y con medicamentos bajo la supervisión e instrucción de su médico. No use este dispositivo en circuitos de anestesia o respiración.

Precauciones referentes al producto LEA LO SIGUIENTE ANTES DEL USO

- Para prevenir choques eléctricos; mantenga la unidad alejada de agua.
- No maneje la unidad o el cable de alimentación con las manos mojadas.
- No sumerja la unidad en líquido.
- No la use al estar tomando un baño.
- No trate de agarrar una unidad que ha caído al agua. Desenchufe la unidad inmediatamente.
- No toque ni intente reparar la unidad.
- No use la unidad si tiene alguna pieza dañada (incluyendo el cable o el conector de alimentación), si ha sido sumergida en agua o si se ha caído. Envíe la unidad sin demora a que se examine y repare.
- La unidad no deberá usarse en un lugar donde se estén usando gas inflamable, oxígeno o productos atomizados.
- Mantenga los orificios de ventilación descubiertos. No coloque la unidad en una superficie blanda en la que pudieran obstruirse los orificios.
- Si la tapa de la medicación está vacía, no trate de operar la unidad.
- Si ocurre cualquier anomalía, no use la unidad hasta que no haya sido examinada y reparada.
- La unidad no deberá dejarse desatendida al estar conectada a la red eléctrica.
- No la incline ni sacuda mientras que está en operación.
- Desconecte la unidad de la toma de corriente eléctrica antes de la limpieza y el llenado, y después de cada uso.
- No use accesorios que no hayan sido autorizados por el fabricante.
- No desmonte o intente arreglar esta unidad
- No utilice este aparato o en circuitos de anestesia o de respiración forzada

ADVERTENCIA: Este símbolo en el producto significa que se trata de un producto electrónico y, en conformidad con la Directiva Europea 2012/19/EU, los productos electrónicos deberán desecharse en su centro de reciclaje local para un tratamiento seguro.

C. Operación de su nebulizador

El dispositivo nebulizador funciona hasta un ángulo de 45°. Si el ángulo supera los 45° no se generará aerosol.

- Conecte un extremo del conector de tubo de aire en la salida de aire.
- Con precaución, conecte el extremo opuesto de conector de tubo de aire en el vástago del dispositivo nebulizador.
- Encaje la boquilla acodada o alguna máscara opcional en la parte superior del dispositivo nebulizador.
- La capacidad de la taza de la medicación es de 2-5 ml.

NOTA: Después de cada uso se recomienda un periodo de reposo de 30 minutos. El compresor se apagará automáticamente si se calienta excesivamente. Si esto sucede, inmediatamente:

- Ponga el interruptor en la posición de APAGADO.
 - Desenchufe el cable de alimentación de la toma de corriente.
 - Permita que el motor se enfríe por 30 minutos.
- Antes de volver a encender la unidad, asegúrese de que los orificios de ventilación no estén obstruidos.
- Verifique la posición del interruptor en la posición de APAGADO.
 - Mantenga sus ojos alejados de la salida de la neblina de medicación.
 - La capacidad máxima de la taza de la medicación es de 5 ml y no deberá sobrepasarse.
 - No use esta unidad al estar conduciendo un vehículo.
 - Si se presenta cualquier malestar o anomalía, pare el uso de la unidad inmediatamente.
 - No utilice este aparato si el tubo de aire está doblado
 - La Pentamidine no es una medicación aprobada para su uso con este aparato

Precauciones de almacenaje

- No guarde la unidad en un lugar expuesto directamente al sol, a altas temperaturas o humedad.
- Mantenga la unidad fuera del alcance de niños pequeños.
- Siempre mantenga la unidad desconectada de la red eléctrica mientras que no la está usando.

Precauciones de limpieza

- Limpiar el filtro de aire, el dispositivo nebulizador, la boquilla y cualquier otra pieza opcional antes de cada uso. Las piezas que estén sucias o desgastadas deberán sustituirse.
- No sumerja la unidad en agua. La unidad podrá quedar dañada.
- Enjuague el dispositivo nebulizador y los componentes con agua caliente de la lavra.
- Sequelas con toallas limpias o permita que se sequen completamente al aire.
- 6. Vuelva a ensamblar el dispositivo nebulizador.
- Elimine siempre cualquier resto de medicación después de cada uso. Use medicación nueva cada vez que utilice este aparato.
- No almacene el tubo de aire con humedad o restos de medicación ya que esto podría ocasionar infecciones bacterianas.

DESCARGO DE RESPONSABILIDAD MÉDICO:

Ni este manual ni el producto se han concebido para sustituir cualquier consejo proporcionado por parte de su doctor u otro profesional médico.

No use la información presentada aquí o otro producto para el diagnóstico o el tratamiento de un problema de salud, o para recetar alguna medicación. Si sospecha tener algún problema médico, consulte a un médico sin demora.

La vida útil de este dispositivo es como sigue, asumiendo que el producto se utiliza para nebulizar 2 ml de medicación unas 2 veces al día durante 8 minutos cada vez a una temperatura ambiente de 25°C. La vida del equipo suministrado puede variar dependiendo de las condiciones ambientales de uso.

Unidad central	5 años
• Nebulizador	1 año
• Tubo de aire ,Boquilla acodada	1 año
• Filtro de aire	60 días
• Máscara de niño, Máscara de adulto	5 años

Especificaciones del producto

Potencia	AC 230V/50Hz o AC 220V/60Hz o AC110V/60Hz
Consumo de potencia	≤ 130 W
Nivel sonoro	≤ 60 dBA (A 1 m de distancia del NE100)
Rango de presión del compresor	≥ 29 psi (200 kPa)
Rango de presión de operación	≥ 15 psi (103 kPa)
Caudal	≥ 3.5 lpm
Rango de temperatura de operación	De 50 a 104°F (de 10 a 40°C)
Rango de humedad de operación	De 10 a 90% RH
Rango de presión ambiental de funcionamiento	700 – 1060 hPa
Rango de temperatura de almacenaje	De -4 a 104°F (de -20 a 60°C)
Rango de humedad de almacenaje	De 10 a 90% RH



DE Deutsch

Einführung

Vielen Dank, dass Sie den Vernebler NE100 mit Kompressor erworben haben. Bei sorgfältiger Verwendung und Pflege wird Ihnen Ihr Vernebler viele Jahre lang zuverlässige Dienste leisten. Dieses Gerät arbeitet mit normaler Wechselspannung. Die Behandlung ist schnell, sicher und bequem, so dass sich dieses Gerät ideal für alle Altersgruppen eignet. Wir empfehlen Ihnen, diese Anleitung gründlich durchzulesen, damit Sie die Funktionen Ihres Verneblers kennen lernen. Der Kompressor für Ihren Vernebler sollte unter Aufsicht eines zugelassenen Arztes bzw. eines Spezialisten für Atemtherapiebehandlungen betrieben werden. Mit Unterstützung Ihres Arztes bzw. Ihres Therapeuten für Atemwegserkrankungen können Sie sicher sein, dass Sie die effektivste Inhalationsbehandlung für Ihre Atemwegserkrankungen erhalten.

HINWEIS: Ihr Vernebler ist zur Behandlung von Asthma, COPD und anderen Erkrankungen der Atemwege vorgesehen, bei denen während der Therapie ein Medikament in Aerosolform benötigt wird. Bitte sprechen Sie mit Ihrem Arzt bzw. Ihrem Apotheker, ob die Ihnen verschriebenen Medikamente für den Einsatz mit dem Vernebler zugelassen sind. Richten Sie sich in Bezug auf Art, Dosis und Medikation nach den Anweisungen Ihres Arztes bzw. Therapeuten für Atemwege.

Dieses Gerät erfüllt die Vorschriften der EU-Richtlinie 93/42/EWG (Medizinäergerechtheit) sowie die Anforderungen der Europäischen Norm EN 13544-1:2007+A1:2009, Geräte zur Atemwegtherapie – Teil 1: Verneblersysteme und ihre Teile.

Bitte lesen Sie die Bedienungsanleitung sorgfältig durch, bevor Sie das Gerät verwenden, und geben Sie diese auf.

Zusammenbau Ihres Inhalators
Folgen Sie den Reinigungsanweisungen in dieser Anleitung im Abschnitt „Reinigungsverfahren“, bevor Sie Ihren Vernebler erstmals bzw. nach längerer Lagerung verwenden.

WICHTIGER HINWEIS: Trennen Sie immer den Kompressor vom Netz und achten Sie darauf, dass der Netzschalter in Stellung "OFF" steht, bevor Sie das Gerät reinigen, zusammenbauen sowie vor und nach jeder Verwendung.

- Stellen Sie den Kompressor auf einer ebenen, stabilen Oberfläche in Reichweite ab.
- Verdrehen Sie den Deckel des Verneblers leicht, und ziehen Sie ihn gerade nach oben ab, um die beiden Teile (Medizinschale und Abdeckung) zu lösen.
- Der Stößel in der Medikamentenschale muss in dem Röhren der Düse sitzen.
- Geben Sie die korrekte Menge des verschriebenen Medikaments in die Medikamentenschale.
- Bauen Sie den Vernebler vorsichtig wieder zusammen, indem Sie Medikamentenschale und Abdeckung gegeneinander verdrehen. Achten Sie darauf, dass die beiden Teile fest sitzen.

Vorsicht

Bitte beachten Sie bei Bedienung Ihres Verneblers die allgemeinen Sicherheitsvorschriften. Dieses Gerät sollte nur für den bestimmungsgemäßen Zweck entsprechend der Beschreibung in dieser Anleitung