Digital Ultra™ User Manual



- Flushmount
- Flexmount
- Digital Newport



INDICATIONS FOR USE

To be used in nitrous oxide-oxygen sedation systems for delivering to a patient a mixture of nitrous oxide and oxygen gases with a maximum nitrous oxide concentration of 70%.

CONTRAINDICATIONS

Contraindications for use of nitrous oxide/oxygen inhalation may include:

- 1. some chronic obstructive pulmonary diseases;
- 2. severe emotional disturbances or drug-related dependencies;
- 3. first trimester of pregnancy;
- 4. treatment with bleomycin sulfate;
- 5. methylenetetrahydrofolate reductase deficiency.
- 6. severe asthma

Whenever possible, appropriate medical specialists should be consulted before administering analgesic/anxiolytic agents to patients with significant underlying medical conditions (e.g., severe obstructive pulmonary disease, congestive heart failure, sickle cell disease, acute oritis media, recent tympanic membrane graft, acute severe head injury.¹ Operator must review patient history with regard to these health issues.

¹ American Academy of Pediatric Dentistry. Clinical Guidelines: Guidelines on Use of Nitrous Oxide for Pediatric Dental Patients. Pediatr Dent 2009;31(6):148-151. Available at: "http://www.aapd.org/ media/policies.asp". Accessed February 23, 2010.

WARNINGS

To be used only by a professional trained in the use of nitrous oxide, using titration method.

Patient should always be closely monitored during nitrous oxide use. If patient has an adverse reaction, reduce or stop the flow of nitrous oxide as needed. The O_2 flush button can be used to rapidly purge the lines of N_2O . If patient does not show signs of quick recovery, remove nasal hood and treat with pure oxygen from either the O_2 resuscitator fitting or an auxiliary oxygen tank using a demand valve, oxygen assisted manual resuscitator, or equivalent. Call for emergency assistance if rapid response is not achieved.

Do not use this device for the administration of general anesthesia or as part of, or in conjunction with, a general anesthesia administration system.

For use with 100-240 V AC, 50/60 Hz only.

Use only hospital grade power cord connected to an equivalent receptacle marked "Hospital Only" or "Hospital Grade." For emergency power shut-off and isolation, unplug power cord from wall outlet.

Unit is calibrated at the factory to $\pm 5\%$ per industry recommendations.

Verify that the correct gases are being delivered to the correct flowmeter inlets by performing the oxygen failsafe test in section VII.

NIOSH recommends N₂O levels in operator area be kept below 25 PPM.

Administer pure oxygen to patient for at least 5 minutes after nitrous oxide procedure.

CAUTIONS

Federal (U.S.) law restricts this device to sale by or on order of a dentist or physician.

Do not attempt to repair, alter, or calibrate this device. Unauthorized repair, alteration or misuse of this device is likely to adversely affect the performance and will void the warranty.

Safety features contained in this notice should be routinely checked to assure proper function. If any of these safety features are not functioning properly, contact your dealer or Accutron and arrange for the necessary repairs before reusing the machine.

Always use clean, dry medical gases. Introduction of moisture or other contaminants into Accutron analgesia gas machines may result in defective operation.

Users of electronic flowmeters that control % oxygen should be aware that Accutron controls % nitrous oxide.

ELECTROMAGNETIC COMPATIBILITY (EMC)

This equipment passes radiated emissions requirements for CISPR 11 Class B devices and EM field immunity requirements of IEC6100-4-3. Operating other radio frequency devices in the vicinity of this equipment should be evaluated accordingly. The minimum distance between this equipment and other RF devices is given by the formula below and the table. The minimum distance is determined by the power of the RF device operating in the vicinity of the digital flow meter.

minimum distance: $d = \frac{\sqrt{(Power)}}{0.388}$ meters

<u>Power</u>	<u>Distance (meters)</u>	Distance (feet or inches)
0.01 Watts	0.258	10 inches
0.1 Watts	0.816	2 feet and 8 inches
1 Watts	2.58	8 feet and 6 inches
10 Watts	8.16 26 feet and 9 inches	
over 10 Watts	Operating the digital flow meter within close proximity of such a transmitter is not recommended.	

Warning: Changes or modifications to the equipment may result in unsafe operation.



Recycling

Old electrical and electronic equipment must be disposed separately and may not be included in the regular domestic waste. Dispose in accordance with prevailing ordinances. Alternatively, the unit can be handed over to Accutron for correct recycling.

INDICATIONS D'UTILISATION

Vous ne devez utiliser cette machine que dans des systèmes de sédation oxygène-oxyde nitreux qui livrent au patient un mélange de gaz d'oxyde nitreux et d'oxygène ayant une concentration maximale d'oxyde nitreux de 70 %.

CONTRE-INDICATIONS

Les contre-indications sur l'inhalation d'oxyde azoteux/oxygène peuvent inclure:

- 1. certaines maladies respiratoires obstructives chroniques;
- 2. des troubles émotionnels graves ou des dépendances liées à la toxicomanie;
- 3. le premier trimestre d'une grossesse;
- 4. le traitement au sulfate bléomycinique;
- 5. une déficience en réductase d'acide tétrahydrofoliqueméthylène.
- 6. sever asthme

Dans la mesure du possible, consulter un spécialiste médical avant d'administrer des agents analgésiques/anxiolytiques aux patients qui présentent des troubles médicaux sous-jacents (par ex.: maladies respiratoires obstructives graves, défaillance cardiaque, drépanocytose, otite moyenne aiguë, greffon récent de la membrane du tympan, traumatisme crânien grave aigu.¹ L'opérateur doit examiner l'historique du patient à l'égard de ces problèmes de santé.

¹ American Academy of Pediatric Dentistry. Lignes directrices cliniques: Lignes directrices sur l'utilisation de l'oxyde azoteux pour les patients dentaires pédiatriques. Pediatr Dent 2009;31(6): 148-151. Disponible sur le site Web: "http://www.aapd.org/media/policies.asp". Accès le 23 février 2010.

AVERTISSEMENTS

Pour être utilisé uniquement par un professionnel formé à l'utilisation de l'oxyde nitreux en utilisant la méthode de titration.

Le patient doit toujours faire l'objet d'une surveillance étroite pendant l'utilisation de l'acide azoteux. Si le patient présente des effets indésirables, réduire ou arrêter le débit de l'acide azoteux, selon le besoin. Le bouton de purge O_2 pour peut être utilisé pour évacuer rapidement les lignes de N_2O . Si le patient n'affiche aucun signe de récupération rapide, enlever l'enceinte nasale de hood et le traiter avec de l'oxygène pur, soit à partir du raccord du réanimateur O_2 ou d'une bouteille d'oxygène auxiliaire utilisant un détendeur, d'un réanimateur manuel assisté par de l'oxygène ou l'équivalent. Demander de l'aide d'urgence si une réponse rapide n'est pas obtenue.

Ne pas utiliser ce dispositif pour l'administration d'une anesthésie générale ou comme partie ou encore en conjonction avec un système d'administration d'anesthésie générale.

Pour une utilisation à 100-240 V AC, 50/60 Hz seulement.

Utilisez le cordon d'alimentation de qualité que l'hôpital connecté à une prise équivalente portant la mention «Hôpital seulement» ou «Hospital Grade». Pour pouvoir d'arrêt d'urgence et d'isolement, débranchez le cordon d'alimentation de la prise murale.

Unité est calibré à l'usine de ± 5% selon les recommandations de l'industrie.

Vérifiez que le bon gaz sont livrés au bon débitmètre anses en effectuant l'oxygène Failsafe Test dans la section VII.

Le NIOSH (USA) recommande $\mathrm{N}_2\mathrm{O}$ niveaux de zone opérateur être maintenu en dessous de 25 ppm.

Administrer oxygène pur pour au moins 5 minutes après utilisation d'azote.

ATTENTION

Conformément à la loi fédérale des États-Unis, cette machine ne peut être vendue que par un dentiste ou par un médecin ou sur leur ordonnance.

N'essayez jamais de réparer, de modifier ou de calibrer cet instrument. Toute réparation, modification non autorisée ou mauvais usage de cet instrument nuira considérablement à son rendement et annulera la garantie.

Vous devez vérifier couramment les caractéristiques de sécurité de cette brochure pour assurer un bon fonctionnement. Si l'une des caractéristiques de sécurité ne fonctionne pas bien, appelez votre revendeur ou Accutron et prenez les mesures nécessaires pour réparer votre appareil avant de l'utiliser à nouveau.

Utilisez toujours des gaz médicaux propres, secs. Toute introduction d'humidité ou d'autre contaminant dans un appareil d'analgésie à gaz Accutron provoquera un mauvais fonctionnement.

Les utilisateurs de débitmètres électroniques qui contrôlent $\%O_2$ doivent être conscients que Accutron contrôle $\%N_2O$.

Compatibilité électromagnétique (CEM)

Cet équipement passe exigences sur les émissions de CISPR 11 Classe B dispositifs et exigences d'immunité de champ EM de IEC6100-4-3. l'autre opération les dispositifs de radiofréquence dans le voisinage de ces équipements doivent être évalués en conséquence. La distance minimale entre cet équipement et d'autres RF appareils est donnée par la formule ci-dessous et le tableau. La distance minimum est déterminée par la puissance du dispositif RF fonctionnant dans le voisinage de la numérique débitmètre.

distance minimale:
$$d = \frac{\sqrt{(Puissance)}}{0.388}$$
 mètres

<u>Puissance</u>	Distance (mètres)	Distance (pieds ou en pouces)
0,01 Watts	0.258	10 pouces
0,1 Watts	0.816	2 pieds et 8 pouces
1 Watts	2.58	8 pieds et 6 pouces
10 Watts	8.16 26 pieds et 9 pouces	
plus de 10 Watts	de fonctionnement du débitmètre numérique dans près la proximité d'un tel émetteur est déconseillée.	

Attention: Les modifications apportées à l'équipement peuvent entraîner des fonctionnement dangereux.



Recyclage

Les anciens équipements électriques et électroniques doivent être éliminés séparément et ne peuvent pas être inclus dans les déchets domestiques réguliers. Éliminer conformément aux règlements en vigueur. Autrement, l'appareil peut être remis à Accutron pour le recyclage correct.

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I. TECHNICAL SPECIFICATIONS

Physical

Flushmount Main Unit 8x4x7.5" (20,3x10,2x19 cm) Weight: 5.2 lbs (2,4 kg) Flushmount Controller 9x4x1.5" (22,9x10,2x3,8 cm) Weight: 1.0 lbs (0,8 kg) Flexmount 8x4x7.5" (20,3x10,2x19 cm) Weight: 8.2 lbs (3,7 kg) Digital Newport 19x32x22" (48x80x56 cm) Weight: 75 lbs (34 kg)

Electrical

100-240 V AC 50/60 Hz. 0.5 A

Socket is IEC/EN 60320-1 C14 and accepts an IEC/EN 60320-1 C13 connector Power cords available for most countries

Internal Battery

Lithium coin battery, 3v CR2032

Fuses

T500mAL, 250v (quantity 2) Accutron part number 26456-FRU

Gas Supply

Oxygen: 50-55 PSI at 100 LPM min. Nitrous Oxide: 50-55 PSI at 10 LPM min.

Gas Fittings

Oxygen Inlet: Male DISS CGA 1240

Nitrous Oxide Inlet: Male DISS CGA 1040

Mixed Gas Outlet: Male DISS CGA 1160 or 1/4" Hose Barb Oxygen Resuscitator Connection: 1/4" I.D. Quick Disconnect

Adapter hoses available for AGA, DIN, NIST, BOC, AFNOR, SIS, UNIFO

Gas Delivery

Oxygen Flush: Minimum 20 LPM

Oxygen Flow: 1.0 LPM-9.9 LPM (Accuracy +/- 0.5 LPM)

Oxygen %: 30%-100%

Oxygen Resuscitator Flow: Minimum 100 LPM, 100% O₂ Nitrous Oxide Flow: 0-6.9 LPM (accuracy ⁺/- 0.5 LPM)

Nitrous Oxide %: 0-70% Mixed Gas Flow: 1.0-9.9 LPM

Environmental

Transportation/Storage: Temperature: -40°C to +70°C (-40°F to 158°F)

Humidity: 10-100%

Operating Conditions: Temperature: 15°C to 32°C (68°F to 90°F)

Humidity: 30-75%

Atmospheric Pressure: 72-101 kPa (543-760 mmHg)

Accessories

Printer

Remote Control

Remote Flow System (RFS)

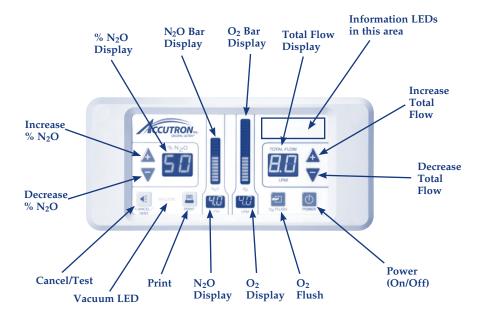
Accu-Vac

Applied Part (rated BF)

Scavenging Circuit II	PN32007
Scavenging Circuit II (RFS)	PN32009
Scavenging Circuit II, ClearView	PN43003
Scavenging Circuit IL ClearView (RFS)	PN43004

II. CONTROL/DISPLAY FEATURES

Front Control/Display Panel



Remote Control



Front Control/Display Panel & Remote Control

<u>Control/Display Panel</u> – Easy-to-use control panel allows setting Total Flow and N_2O concentration with the touch of a finger. Bright, color-coded displays make it simple to monitor status, even from a distance. Optional hand-held Remote allows functional control at patient chairside.

<u>Total Flow Control</u> – Allows setting of Total Flow in either (.1) LPM or (.5) LPM increments. Once set, the Total Flow will remain constant even if % N₂O is changed.

<u>Total Flow Display</u> – Shows Total Flow in numeric LPM format with yellow LEDs.

 $\frac{\% \text{ N}_2\text{O Control}}{\text{Control}}$ – Allows setting of N₂O concentration in either 1% or 5% increments up to a maximum of 70%. Once set, % N₂O will remain constant even if Total Flow is adjusted.

% N₂O Display – Shows % N₂O in numeric format with blue LEDs.

<u>Power</u> – Turns unit on and off with the push of a button.

<u>Flush Control</u> – Ceases N_2O Flow and provides a minimum of 20 LPM of pure O_2 as long as button is pressed.

 O_2 Flow – Two green LED displays show O_2 Flow in both LPM numbers and graphic bars simulating standard flow tube.

 N_2O Flow – Two blue LED displays show N_2O Flow in both LPM numbers and graphic bars simulating standard flow tube.

<u>Print</u> – When optional printer is installed, provides printout of current conditions.

<u>Cancel/Test</u> – During an alarm mode, turns off audio alarm for one minute. During normal mode, performs self-check of all displays.

<u>Information LEDs</u> – Informs administrator of failures with O_2 supply, N_2O supply, O_2 flush, or mixed gas line to patient. Also indicates if remote vacuum is connected.

III. FLOWMETER MODELS

FLUSHMOUNT MODEL



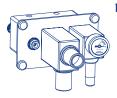
Two piece design for easy installation into a cabinet or a wall. Thin control module located away from main module, power, and gas lines, requires remote outlet.

Flushmount Remote Outlets Options



a) Standard

(Contains mixed gas outlet and O₂ resuscitator connection)



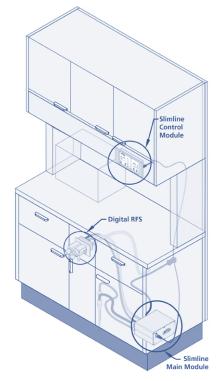
b) Remote Flow System (RFS)

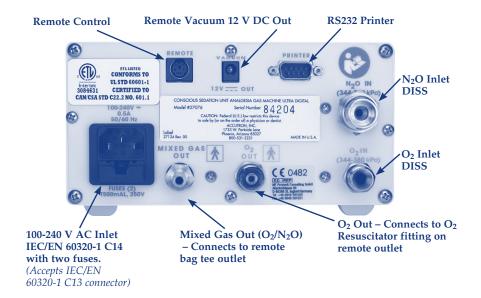
(Contains mixed gas outlet, O₂ resuscitator, and vacuum controller with gauge display)



(Contains mixed gas outlet, O₂ resuscitator, and vacuum controller with LED display)

Sample Installation





Flushmount Main Module Utility Connections

 O_2 Inlet – Labeled O_2 IN – Equipped with male CGA 1240 DISS fitting to prevent gas mix-up.

 $\underline{\text{N}_2\text{O}}$ Inlet – Labeled N₂O IN – Equipped with male CGA 1040 DISS fitting to prevent gas mix-up.

 $\underline{O_2}$ Out – Labeled O_2 OUT – Equipped with male CGA 1240 DISS fitting, provides oxygen for resuscitator.

Mixed Gas Port – Equipped with male CGA 1160 DISS fitting (Medical Air)

<u>Power Cord Inlet</u> – Clearly labeled 100-240 V AC, 50/60 Hz and protected by two fuses for added safety. Fuses are 240 V, 0.5A, slow-blow. To disconnect power, unplug cord. Do not position where this disconnection would be difficult.

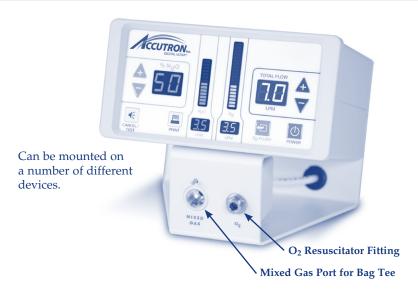
<u>RS232 Printer Connector</u> – Clearly labeled and designed for use with optional Accutron-approved printer.

<u>Remote Control Connector</u> – Remote Control connection. Using the Remote Control, general adjustments can be made from a distance of up to 10 feet away from device.

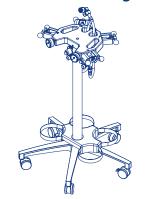
<u>Remote Vacuum Valve Connector</u> – Clearly labeled and designed to actuate a vacuum valve in the Accu-Vac Automatic Vacuum Scavenging Port (PN 56955).

 $\underline{O_2}$ Out Connector – Direct connection to oxygen for use with resuscitator. Provides minimum of 100 LPM of oxygen.

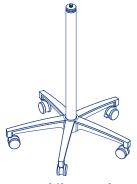
FLEXMOUNT MODEL



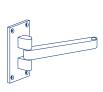
Flexmount Mounting Options



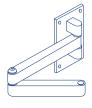
4-Cylinder Portable System



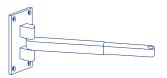
Mobile Stand



9" Wall Arm

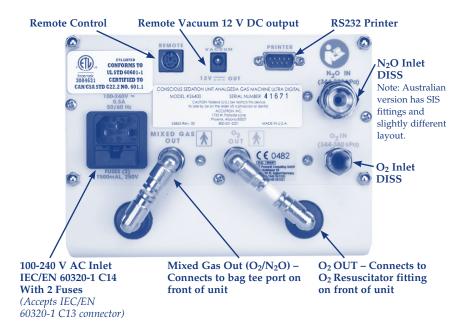


16" Bi-fold Wall Arm

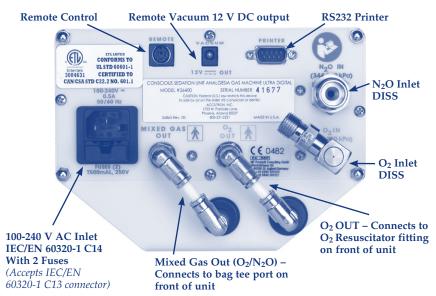


Telescoping Wall Arm (Extends 12"-17")

Wall Arm and Mobile Stand version



4 Cylinder version



Flexmount Utility Connections

 O_2 Inlet – Labeled O_2 IN – Equipped with male CGA 1240 DISS fitting to prevent gas mix-up.

 $\underline{N_2O}$ Inlet – Labeled N_2O IN – Equipped with male CGA 1040 DISS fitting to prevent gas mix-up.

O2 Out – Labeled O2 OUT – Connects to oxygen resuscitator fitting on front.

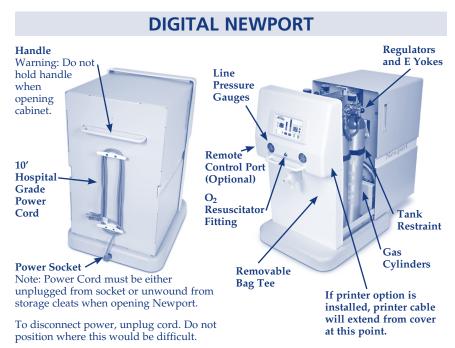
Mixed Gas Out - Connects to bag tee port on front.

<u>Power Cord Inlet</u> – Clearly labeled 100-240 V AC, 50/60 Hz and protected by two fuses for added safety. Fuse ratings are 240 V, 0.5 A, slow-blow. To disconnect power, unplug cord. Do not position meter where this disconnection would be difficult.

<u>RS232 Printer Connector</u> – Clearly labeled and designed for use with optional Accutron-approved printer.

Remote Control Connector – Remote Control connection. Using the Remote Control, general adjustments can be made from a distance of up to 10 feet away from device.

O₂ Out Connector – Direct connection to oxygen resuscitator fitting on front. Provides minimum of 100 LPM of oxygen.



Digital Newport Setup and Connections

<u>View</u> – When looking at the front of the Newport cabinet, Nitrous Oxide is on the left as shown by the blue Nitrous Oxide gauge. Oxygen is on the right as shown by the oxygen gauge.

<u>Opening Newport Cabinet</u> – Open the Newport cabinet by placing one hand on the front cover and the other in the handle on the side. Pull handle back all the way. Make sure the power cord is unplugged from the socket in the skirt first.

Install Gas Cylinders -

- 1. Undo Velcro straps
- 2. Angle tank into position
- 3. Strap tanks loosely
- 4. Remove regulators from accessories box
- 5. Attach regulators to tanks by loosening yoke handle, verifying seal is in place, aligning seal with gas opening, aligning pins with holes in yoke, tightening yoke onto cylinder

NOTE: Regulators are pinned to match gas cylinders. Connect gas hoses to regulator outputs.

NOTE: Hoses and regulators use DISS fittings to prevent mix up. Rotate cylinders so that regulators fit inside.

<u>Open Gas Cylinders</u> – Pick one tank of each gas to be "in use" tank and attach "in use" label to cylinder. Open these cylinders using the cylinder wrench provided in accessory box. Close Newport cabinet.

Connect Power -

- 1. Plug cord back into skirt outlet
- 2. Plug other end of cord into receptacle

NOTE: System will operate from 100-240 V, 50-60 Hz. Internal power supply will switch automatically.

Scavenging Circuit -

- Install bag tee from accessories box into front of Newport.
- Attach scavenging circuit to bag tee.
- Connect vacuum hose to system vacuum.
- Install nasal hood into scavenging circuit and start vacuum flow per scavenging instructions.

Start up - See Section VII, Directions For Use

Digital Newport: Managing Gas Supply

Note: Power Cord must be either unplugged from socket or unwound from storage cleats when opening Newport.

- 1. Identify the two in-use tanks (N₂O and O₂) by attaching "in-use" identification tags to them.
- When one of the in-use tanks is empty, close the tank by turning the valve clockwise. Leave the empty tank in place.
- 3. Move the "in-use" tag to the appropriate reserve tank, which now becomes the in-use tank. Replace the empty tank with a new tank, which then becomes the reserve tank.
- 4. Open the valve of the new in-use tank by turning counterclockwise.
 - Note: Do not open the reserve tank until the in-use tank is empty. When changing gas tanks, make certain both same gas cylinders that are involved in the exchange are closed. Always remember to switch the "in- use" tags over to the new tanks.
- 5. Contact gas supplier to request new gas cylinders as needed.

Digital Newport: Changing Gas Cylinders

Always use clean, dry medical grade gases. Introduction of moisture or other contaminants into Accutron Analgesia gas machines may result in defective operation.

The Digital Newport Flowmeter System is designed to operate with a 50-55 psi line pressure on each gas (O_2 and N_2O). The line pressure can be determined by reading the O_2 and the N_2O line pressure gauges that are located on the front panel of the Newport.

The Newport has features incorporated to make cylinder change-out easy. To assure proper operation of the system, 4 cylinders must be properly installed at all times. Please review the following cylinder replacement procedures before attempting to change cylinders.

Note: Power Cord must be either unplugged from socket or unwound from storage cleats when opening Newport.

- 1. Open the tank enclosure by sliding the cover to the rear.
- 2. Close the valve on the empty tank (clockwise).
- 3. Release the Velcro restraint on the empty tank.
- 4. Tilt the empty tank out slightly.
- 5. Loosen the round handle on the regulator yoke.
- 6. Remove the regulator from the empty tank (make certain that the regulator is not dropped or damaged while it is off the tank).
- 7. Lay the empty cylinder down on the floor (cylinders should only stand upright when properly restrained as they can easily fall over).
- 8. Pick up the full replacement tank and place the bottom of the tank onto the floor of the Newport, leaving it slightly tilted outward for easy replacement of the regulator.
- 9. Check to assure that the sealing gasket has remained on the regulator's E yoke.
- 10. Reattach the regulator to the new full tank by sliding the yoke (attached to the regulator) over the top of the tank post aligning the index pins with the corresponding holes in the tank post.
- 11. Tighten the round handle on the regulator yoke.
- 12. Push the tank into an upright position. Attach and tighten the Velcro tank restraint.

Digital Newport: Control Module and Main Module Locations





• Main flowmeter module is behind shield at rear of unit. Fuses are located in fuse tray at power cord socket on back of flowmeter.

IV. SAFETY FEATURES

 O_2 Fail Safe System – Offers assurance that N_2O ceases if the O_2 supply is interrupted or reduced. Alarm will sound, FAILURE OXYGEN LEDs will light, and N_2O flow will cease.

 N_2O Supply Warning – Triggers an alarm if the N_2O supply cannot keep up with user's needs. Alarm will sound and FAILURE NITROUS LEDS will light. Flow switchs to 100% oxygen.

<u>Mixed Gas Flow to Patient</u> – Monitors pressure in mixed gas line and triggers an alarm if flow to patient is obstructed. Alarm will sound and FAILURE NITROUS OXYGEN LEDS will light.

<u>Emergency Air Valve</u> – Automatically provides patient with ambient air if gas flow is interrupted. Located on remote bag tee.

<u>Directional Check Valve</u> – Prevents re-breathing of exhaled gases and protects against CO₂ buildup. Located on remote bag tee.

 O_2 Resuscitator Connection – Direct connection to O_2 for use with resuscitator. Port is always open and provides 100 LPM minimum of pure oxygen.

V. GENERAL INSTRUCTIONS

1. Warranty

Unpack Digital Ultra and inspect to make certain that the unit has not been damaged during shipment.

The unit's serial number is located on the back of Digital Ultra (see page 17 for Digital Newport's serial number location). Record the serial number in the area provided at the back of this booklet. Once Digital Ultra Flushmount is mounted into a cabinet, the serial number will no longer be visible. Always reference the serial number when corresponding with Accutron.

Complete the warranty card supplied with Digital Ultra and mail to Accutron. Completion of this step ensures proper device warranty coverage. Accutron Digital Ultra analgesia machines carry a two-year limited warranty (see Warranty on page 26 for details).

2. Gas Supply Connection

Always use clean, dry medical gases. Introduction of moisture or other contaminants into Accutron analgesia gas machines may result in defective operation.

After device installation, connect the oxygen and nitrous oxide supply lines to the Diameter Specific Instant fittings or DISS fittings located on the back of the device. See pages 14-16 for information specific to the Digital Newport. It is important that the regulators for both gases be set to give pressures in the range of 50-55 PSI.

Verify that the correct gases are being delivered to the correct ports on device before initial use. This can be done by shutting off the O_2 cylinders and turning the unit on. The unit should immediately go into an oxygen failure fault mode. If unit indicates O_2 gas is flowing, lines are crossed. Correct before using.

3. Power Connection

Device is designed to run on 100-240 V AC and 50/60 Hz. There is no need to flip an external switch; the unit automatically adjusts for any power. A medical-grade power cord is supplied with the unit and must be used to ensure proper grounding. It is important that the power cord be plugged into an outlet marked "Hospital Only" or "Hospital Grade." The power cord is plugged into the AC entry, which contains two fuses and is located on the back of the device. International power cords are offered for most countries. Use only 240 V, 0.5A, slow-blow fuses. Fuses can be ordered from Accutron as part number 26456-FRU.

4. Auxiliary Connections

Plug remote control into remote port on back of Digital Ultra or on front cover of Newport if ordered as option.

Plug Accu-Vac signal line into vacuum connector on back of Digital Ultra (Flushmount only). Vacuum LED should light when Digital Ultra is powered indicating vacuum is flowing.

Plug printer cable from Able printer, if used, into RS232 printer port. Do not use this port for any other devices. If printer is ordered on Newport as option, printer cable will extend from front cover. Note: Use only Able printer supplied by Accutron.

VI. Directions For Use

Read instructions completely before operating Digital Ultra device.

Note: The steps listed below provide a basic functional description of the Digital Ultra usage. A training course that emphasizes a practical, hands-on approach combined with instructions on safe techniques for administration of nitrous oxide-oxygen conscious sedation is recommended before use of this flowmeter. Do not use this device for the administration of general anesthesia or as part of, or in conjunction with, a general anesthesia administration system. Unit must be connected to O_2 and O_2 0 gases at 50-55 PSI. Unit must be plugged into 100-240 V AC, 50/60 Hz power outlet. Use titration method for administering nitrous oxide.

Power button will shut off Digital Ultra normal functions (reference diagram on page 8). For Emergency shutoff, unplug power cord.

NOTE: Detailed information on users, patients, and conditions can be found at the end of section VI.

General Operating Instructions

Setup:

Verify the presence of separate means of oxygen resuscitation and that appropriate staff are trained in its operation.

WARNING: In cases of adverse reactions, discontinue nitrous oxide procedure, remove scavenging circuit, utilize auxiliary resuscitation measures and equipment, and contact Emergency Responders (911).

Verify that oxygen and nitrous gases have not been mixed up by performing the Oxygen Failsafe Test on page 23. Note: the only significant source of injury from nitrous oxide procedures is gas mix-up.

Open inlet gas supplies for oxygen and nitrous oxide by:

- a. Turning on manifolds in central systems or
- b. Opening gas cylinders in Newports and 4 cylinder mobile stands

Verify scavenging vacuum is turned on and properly set per scavenging circuit Instruction for Use

Verify power to flowmeter

Operation:

- 1) <u>Push POWER button</u> on front membrane or on Remote Controller. (Initial flow is set to 5.0 lpm at factory but can be adjusted per instructions on page 21)
- 2) <u>Install scavenging circuit</u> per scavenging circuit Instructions for Use.
- 3) Adjust proper breathing flow by using the TOTAL FLOW and TOTAL FLOW buttons. Reservoir bag should inflate and deflate in response to patient breathing but never go completely full or completely empty. (Increments are set at factory to 0.5 lpm but can be reset to 0.1 lpm increments per instructions on page 20).
- 4) <u>Gradually add nitrous oxide</u> by using the % N₂O. It is recommended that % start at 10% and be increased by 10% intervals until desired reaction is achieved. Allow time between each change for patient to adjust since rapid increasing of N₂O % can cause nausea. (Increments are set at factory to 5% but can be adjusted to 1% per instructions below)

Note: Total flow does not change if % N₂O changes and % N₂O does not change if Total Flow changes.

<u>Caution!</u> Accutron controls % N₂O. Refer to page 22 for possible Use Error for those who are used to meters that control % O₂.

- 5) Max % N₂O is set at the factory to 70%. This limit can be changed to 50% by following instructions on page 21.
- 6) Adjust Total Flow and % N₂O as needed to maintain proper anxiety relief and proper flow levels.
- 7) Turn % N₂O to 0 at the end of the procedure by using the % N₂O. Continue with pure oxygen for 5 minutes.
- 8) Stop gas flow by pushing POWER at the end of the procedure. Remove scavenging circuit and close vacuum.

Additional Functions:

<u>Print</u> - If a Printer is installed, a snap shot of current O_2 flow, N_2O flow, Total Flow, % N_2O , Time and Date can be printed by pushing PRINT. <u>Mute</u> - During an alarm condition, sound is muted by pushing CANCEL/TEST Led Test - During normal operation, CANCEL/TEST will cycle though all Leds to verify all are operational <u>Flush</u> - A minimum 20 lpm of pure oxygen will flow when FLUSH is pushed

Setting Step Size of Total Flow and % N₂O

- 1. To set the % N₂O step size to 5%: Plug in Digital Ultra and leave off. Press and hold % N₂O Up (+) button until beeps (approx. 5 seconds).
- 2. To set the % N₂O step size to 1%: Plug in Digital Ultra and leave off. Press and hold % N₂O Down (-) button until beeps (approx. 5 seconds).
- 3. To set Total Flow step size to 0.5 LPM increments: Plug in and turn off Digital Ultra. Press and hold Total Flow Up (+) button until beeps (approx. 5 seconds).
- 4. To set Total Flow step size to 0.1 LPM increments: Plug in and turn off Digital Ultra. Press and hold Total Flow Down (-) button until beeps (approx. 5 seconds).

Setting Date and Time

- 1. Plug in Digital Ultra and leave off.
- 2. Press and hold Print button down until beeps (approx. 5 seconds). This will start year setting.
- 3. Display will show 20 in % N₂O box and flashing display in Total Flow box.
- Adjust to current year using Total Flow Up/Down buttons. Push % N₂O Up
 (+) button to store value and proceed to next step.

- 5. % N₂O box now displays months. Set the month with Total Flow Up/Down buttons. Push % N₂O Up (+) button to store value and proceed to next step.
- 6. Total Flow box now displays days. Set the day with Total Flow Up/Down buttons. Push % N₂O Up (+) button to store value and proceed to next step.
- 7. N_2O box now displays the hours. Set the hour with Total Flow Up/Down buttons. Push % N_2O Up (+) button to store value and proceed to next step.
- 8. N₂O box now displays minutes. Set the minute with Total Flow Up/Down buttons. Push the cancel button to save value and exit program.

Changing Starting Total Flow Rate

With power connected and flowmeter off, press and hold CANCEL/TEST and Total Flow + (up) Arrow together until beep (5-7 seconds). Current Total Flow will be shown. Adjust to desired flow rate using Total Flow + Up and Total Flow - Down buttons. Acceptable values are 1.0 to 9.9 LPM. Press Flush button to accept new value. Starting Total Flow will now start at the new value.

Changing Max N₂O Percentage (50% 70%)

With power connected and flowmeter off, press and hold CANCEL/TEST and N_2O % Up + buttons together until beep (5-7 seconds). This will toggle between 50% and 70%.

Changing Operating Mode (Normal/Demo)

With power connected and flowmeter off, press and hold CANCEL/TEST and $\rm O_2$ FLUSH buttons together until beep (5-7 seconds). This toggles between Normal (gas flow) mode and DEMO (no gas flow). When in Demo mode, unit will start up flashing SALES for 5 seconds. In this mode, controls and displays will act normal but no gas will be needed or allowed to flow.

Set to Constant 50/50 Mode

This is done at the factory via an internal switch. In 50/50 mode, unit will supply a constant 50% Nitrous Oxide/50% Oxygen mix. Total Flow can be adjusted but will remain 50/50. As a safety feature, the unit can be toggled back and forth between 50/50 and 100% oxygen by using the Nitrous % Up and Down Arrows. 50/50 mode is often used in ERs to minimize shock and in Obstetrics for labor pains.

Application Specifications

Intended Use

The Accutron Digital Flowmeter is used to reduce anxiety in dental patients so that traditional treatments can be performed more efficiently and with less stress on the patient.

Intended Medical Indication

Nitrous oxide in Accutron approved applications is used for anxiety relief and is not a treatment for any particular medical indication per se. It is used to calm patients so that other treatments may be performed. It is proscribed for conscious sedation only and not for use with other anesthesia agents.

Intended patient population

Nitrous oxide is routinely given to healthy individuals over 2 years of age without distinction of age, sex, or weight. See pages 2 and 3 for contra-indications.

NOTE: Patient reaction to nitrous oxide can vary widely between individuals and even in same individuals over time. Some individuals can not tolerate any N₂O.

Intended part of the body

Nitrous oxide is a gas and is delivered to the nose via a nasal hood connected to gas supply tubing. The patient inhales the gas which is then absorbed in the lungs.

Intended user profile

Medical professionals such as doctors, dentists, dental assistants, dental hygienists, and nurses who have been formally trained in the use of nitrous oxide and who have studied the operating instructions in the User Manual.

Intended conditions of use

The Accutron Digital flowmeter is intended to be used in medical facilities such as dental offices and hospitals with standard controlled room environments. See section I for further environmental specifications. Units may be mounted in cabinets, on extension arms, on sliding brackets, or on mobile carts. Units are not meant to be hand mobile by personnel and are not meant to be used in emergency vehicles. Frequency of use can vary from rare (once a month) to (frequent) 10-15 times per day. The user interface is designed for asepsis cleaning.

Description and operating principle

The Accutron Digital PC Flowmeter is a microprocessor-controlled device that precisely meters Oxygen and Nitrous Oxide gases for analgesic purposes in Dental Offices and Hospitals. The flowmeter is powered by 100-240 VAC, 50/60Hz via a power cord that is plugged into a filtered inlet. This voltage is then automatically converted to 24 VDC by a switching medical grade power supply. All components within the flowmeter as well as the user interface are operated on 24 VDC or less.

Precise flow of both oxygen and nitrous oxide are achieved through a feedback circuit containing a proportioning solenoid, a medical grade mass flow sensor, and the main microprocessor. The mass flow sensors measure flow electronically giving real time updates to the circuit board which directs the solenoids to maintain desired flow rates. The processor monitors these flows and gives alarms as needed if the flow of either gas can not match the desired rates. The software also prevents nitrous oxide from flowing without oxygen and maintains oxygen levels at 30% or more at all times.

Clinical operation is very simple as there are only two items to control, TOTAL FLOW and $\%N_2O$. Press POWER to start flow which comes on at 5.0 LPM Total Flow and 0 $\%N_2O$. Adjust Total Flow with the + and – keys until it matches the patients breathing rate. Increase % N_2O by using the + and – keys until proper relaxation is achieved. Decrease % N_2O to 0 at end of procedure and continue with pure oxygen for 5 minutes. Turn Power off. Total Flow and $\%N_2O$ can be adjusted during procedure as needed. Pure oxygen flush is available at any time. Print option is also available.

Possible Use Errors

- Administering too much N₂O or increasing the %N₂O too quickly
- Mistaking the %N₂O with %O₂ used by other electronic flowmeters
- Not taking sufficient patient history before starting procedure
- Not reading the instructions for use or undergoing nitrous oxide training prior to starting procedure

Abnormal Use

Disregarding the instructions and warnings is considered abnormal use.

Training

The intended use is the basis for training which is considered mandatory for all operators prior to use. General Theory on nitrous oxide use and procedures are described in such books as *Nitrous Oxide and Oxygen Sedation* by Clark and Brunick and *Lachgas* by Dr Mathers. It is recommended that users undergo classroom training as well as hands on lab instruction. US training can be found in numerous ADA sponsored continuing education courses. European training can be obtained through the Institute for Dental Sedation, Cologne, Germany (http://www.sedierung.com) or through BIEWER medical (http://www.biewer-medical.com/en/sedation-training), Koblenz, Germany. Consideration of currently valid local and national guidelines for training is required.

VII. PERIODIC EQUIPMENT CHECKS

CAUTIONS

Do not attempt to repair, alter, or calibrate this device. Unauthorized repair, alteration or misuse of this device may adversely affect the performance and will void the warranty.

IMPORTANT

Safety features contained in this notice should be routinely checked to assure proper function. If any of these safety features are not functioning properly, contact your dealer or Accutron and arrange for the necessary repairs before reusing the machine.

Display Test

Check Before Each Use

Turn Digital Ultra on and press CANCEL/TEST button. All displays should light sequentially with an audio beep, including alarms.

<u>Cleaning</u>

Perform After Each Use

Wipe with damp cloth. Do not allow liquids to touch rear auxiliary connections or enter case. Do not spray directly onto unit.

Oxygen Flush Valve Test

Check Monthly

- Turn Digital Ultra on at 5.0 LPM and leave nitrous oxide at 0%.
- Disconnect the corrugated tube from the bag tee outspout. Reservoir bag should remain connected to the bag tee downspout.
- Push the Flush button on Front Panel or Remote Control while blocking the flow from the outspout.

Reservoir bag should fill within 5-10 seconds. Remove any blockage when test is complete.

Outspout Check Valve Test

Check Monthly

- Digital Ultra should be off.
- Reservoir bag should be connected to the bag tee downspout.
- Mixed Gas tube should be connected to the outspout but not connected to the white patient delivery tubing.
- Breathe into the open end of the corrugated tube.

The reservoir bag should not fill.

Override Air Valve Test

Check Monthly

- Digital Ultra should be off.
- Reservoir bag should be connected to the bag tee downspout.
- Mixed gas tube should be connected to the outspout but not connected to the white patient delivery tubing. Reservoir bag should be deflated.
- Draw air through the open end of the corrugated tube.

The Override Valve should open and allow air to enter the corrugated tube. Place your finger over the Override Valve and remove it to verify that air is entering the bag tee through the Override Valve. If the Override is not functioning, do not use Digital Ultra until it is repaired or replaced.

Factory Check

Recommended every 2 years or more frequently as required by prevailing local and national codes.

Send equipment to authorized Service Center for operational check, battery replacement, and recalibration. Note: battery used for time and date info only, loss of battery does not affect safety or operation.

Routine Maintenance

There is no required routine maintenance as long as recommended factory check is at least in the recommended period or more frequently as required by prevailing local and national codes. Unit should only be repaired by trained individuals with specialized equipment. Unauthorized opening of equipment will put individual at risk of electrical shock and will void Warranty.

Disinfecting

The following products are approved for disinfecting the front touch pad area. <u>Do not allow liquid to enter any connectors on back of unit.</u> Do not spray directly onto unit.

Lysol® Brand II I.C. Disinfectant Spray EcoTru® Spray ProSpray Surface Disinfectant COE Foam II CaviCide® Microstat2TM NOTE: DO NOT USE BLEACH

VIII. TROUBLESHOOTING GUIDE

PROBLEM	POSSIBLE CAUSE	REMEDY
Display off	Power cord loose No power to wall outlet Fuse blown	 Connect cord firmly Check circuit breaker and/or call electrician Replace fuse in holder under AC socket.
Oxygen Failure alarm	Low O ₂ pressure	 Verify O₂ input to flowmeter Verify valves are open
Nitrous Failure alarm	Low N ₂ O pressure	 Verify pressure in N₂O cylinder Verify valves are open
Nitrous & Oxygen Failure alarm	Obstructed flow to patient	Remove kinks or obstructions from mixed gas line to patient
Front panel buttons are not operating	 Defective panel Internal problem 	Contact Accutron Customer Service
Remote Control is not operating	Loose connection to unit Defective Remote Internal problem	Check remote plug connection on back If problem persists contact Accutron Customer Service
Displays don't work	Broken display/circuit board	Contact Accutron Customer Service
Automatic Vacuum port (Accu-Vac) doesn't work	Loose connection in back Defective valve Internal problem Insufficient vacuum	1. Check plug connection 2. Verify output of vacuum outlet is 12 +/- 3 V DC 3. Verify system vacuum is > 8 in Hg 4. If not solved, contact Accutron Customer Service
Flush does not have enough oxygen flow	Filter clogged O ₂ solenoid damaged	Contact Accutron Customer Service
Flush does not work at all	1. Internal Problem	Contact Accutron Customer Service
Flow doesn't appear to be accurate	Calibration error Internal problem	Check output with Oxygen Analyzer if available Contact Accutron Customer Service
Unit freezes, doesn't respond, gas won't shut-off	1. Processor problem	Unplug power cord Plug back in (Reboot)

IX. WARRANTY

ACCUTRON 2-YEAR FLOWMETER LIMITED WARRANTY

IF AN ACCUTRON FLOWMETER NEEDS TO HAVE REPAIR WORK OR REPLACEMENT PARTS DURING THE 2-YEAR WARRANTY PERIOD DUE TO MANUFACTURING DEFECTS, ACCUTRON WILL PROVIDE THE PARTS AND LABOR AT NO CHARGE. THE FLOWMETER OWNER IS RESPONSIBLE ONLY FOR A \$35.00 SHIPPING AND HANDLING FEE, WHICH WILL BE ASSESSED EACH TIME A FLOWMETER IS RETURNED TO ACCUTRON FOR WARRANTY WORK.

WARRANTY TERMS

Limited Warranty and Disclaimer: Accutron ("Seller") warrants that its product will be free from manufacturing defects subject to the terms, conditions, and limitation set forth hereinafter, for a period of 2 years for flowmeters and 1 year for other equipment products. Seller's obligations under this limited warranty are contingent on Buyer's full payment of the product purchase price. Except as specifically set forth above, Seller and its affiliates make no warranties, expressed or implied, and specifically disclaim any warranties of merchantability or fitness for a particular purpose.

The liability of Seller and its affiliates for any claims, losses, damages, or expenses from any cause whatsoever (including acts or omissions of third parties) regardless of the form of the action, whether in tort, contract, or otherwise, shall not exceed the repair cost, replacement cost, or purchase price of the product that directly gives rise to the claim. Seller and its affiliates shall not be liable for any incidental, special, reliance, consequential, or indirect loss or damage rising out of this agreement or the products. As used in this paragraph, consequential damages include, but are not limited to, lost profits, lost revenues, property damage, personal injury damage to the Purchaser or third parties, loss of business or profits, and/or loss of business reputation. It is the sole responsibility of Purchaser to determine the suitability of the products for the Purchaser's intended use. Seller's obligation to repair, replace or refund, as set forth above shall be Buyer's exclusive remedy.

This warranty constitutes the entire warranty. This warranty and Seller's liability hereunder shall be construed according to the laws of the State of Arizona without regard to conflict of lay principles.

To activate the Flowmeter Warranty, complete and mail the warranty registration card that accompanies Flowmeter.

Accutron warranties are subject to the following conditions: Accutron products and equipment are warranted to be free from defects in material and workmanship under normal use and service, including all component parts. This warranty shall not apply to defects resulting from accidents, alterations, or misuse. If modifications have affected the operation of the product to render it faulty, this warranty shall be void. This warranty shall be void if any part not of Accutron's manufacture or supply has been incorporated into the product.

THIS WARRANTY IS GIVEN IN PLACE OF ALL OTHER WARRANTIES EXPRESSED OR IMPLIED, OF MERCHANTABILITY, FITNESS FOR A PARTICULAR PURPOSE OR OTHERWISE.

No statement or claim about the product by any employee, agent, representative or dealer of Accutron, Inc. shall constitute a warranty by Accutron, Inc. or give rise to any liability or obligation of Accutron, Inc.

X. WARRANTY AND RETURNED GOODS POLICY

All warranty resolution issues and merchandize returns will be handled through the local authorized Accutron distributor. Contact distributor where unit was purchased.

XI. REPAIR SERVICE POLICY

All service issues will be handled through the local authorized Accutron distributor. Contact distributor where unit was purchased. Check Troubleshooting Guide on page 25 prior to contacting distributor.

XII. ASSISTANCE

For Assistance, contact your local dental distributor or call Accutron Customer Service at:

Toll-free: (800) 531-2221 Local: (623) 780-2020 Fax: (623) 780-0444

Hours of operation: 7:00 AM – 4:30 PM MST

Service ship-to address:

Accutron, Inc. 1733 W. Parkside Lane Phoenix, AZ USA 85027

Visit our website:

www.accutron-inc.com



MT Promedt Consulting GmbH Altenhofstrasse 80 D-66386 St. Ingbert/Germany Tel. +49 (0) 6894-581020 Fax +49 (0) 6894 581021



XIII. OWNERSHIP INFORMATION

Dr. Name:	
Street Address:	
City/State/Zip:	
Accutron Digital Ultra Device Serial Number:	



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