

LIFEPAK[®] 1000

Defibrillator



Highly configurable to adapt to your resuscitation protocol

Upgradeable platform to meet your resuscitation needs

The most rugged defibrillator ever built by Medtronic

Backed by our heritage of quality, reliability and service

Logical, natural interface with a large, intuitive screen

The nonrechargeable battery can deliver over 425 shocks

The LIFEPAK 1000 defibrillator is designed to be the most powerful basic life support (BLS) tool available. Fully customizable, it can adapt to the protocols of today and the future. cprMAX™ technology is highly flexible, with the most configurations available to accommodate your patient and CPR protocol requirements as of July 2005. Future patient focused upgrades will help you provide the most advanced care in a way that compliments how you work best. The 1000 is the most rugged defibrillator ever built by Medtronic, with an IP55 rating. ADAPTIV™ biphasic technology provides a full range of energy up to 360 joules.

With a large screen, graphics and text are clear and easy to read at a distance. The LCD screen can provide both graphical and ECG displays. A manual defibrillation override option is also available.

The advanced battery system in the LIFEPAK 1000 defibrillator provides more power for your patients. A visible and accessible battery gauge gives you the ability to assess battery readiness with the quick touch of a button. Our battery also performs advanced communications directly with the device so you always know the battery's condition.

The 1000 is compatible with other Medtronic products and services across the full spectrum of care, from the field to the hospital, including data download.

DEFIBRILLATOR

All specifications are at 20°C unless otherwise specified.

Waveform: Biphasic truncated exponential with voltage and duration compensation for patient impedance*

Energy Sequence: User configurable, 150 joules–360 joules. Default energy output settings are 200, 300, 360 joules. 360 joules for every shock thereafter.


Charge Time: With new, nonrechargeable battery pack; 200 joules in less than 7 seconds (360 joules in less than 12 seconds)

3-wire (Lead II) monitoring capability standard with device: Requires purchase of 3-wire (Lead II) monitoring cable and LIFE-PATCH® electrodes

Device Software: Field upgradeable

Infant/Child Reduced Energy Defibrillation Electrodes: Reduces selected energy by a factor of 4. Intended for use only with children up to 8 years of age or 25kg (55 lbs).

Safety Classification: Internally powered equipment IEC 60601-1/EN 60601-1

Electrical Protection: Input protected against high voltage defibrillator pulses per IEC 60601/EC 60601 

*Specifications apply from 25 to 200 ohms. Voltage compensation is limited to the voltage that would result in delivery of 360 joules into 50 ohms.

DEVICE SETTINGS

Modes:

- **AED** – Provides operating capability for basic users
- **Manual** – Provides operating capability for advanced users
- **ECG** – Provides ECG display capability with 3-wire ECG cable
- **Setup** – Allows user to configure the device
- **Data Transfer** – Allows user to transfer patient data
- **Auto Test** – Provides daily automatic tests of hardware and software

Controls: On/Off, Shock, Menu, Two (2) configurable soft keys

User Defined Options:

- **Device ID** – Assigns unique identifier to particular device
- **Energy Sequence** – User configurable from 150 to 360 joules
- **Flexible Energy** – Increases only after a lower energy was unsuccessful
- **Auto Analyze** – User can configure device to auto analyze, auto analyze after first shock, or prompt user to push analyze key before each analysis period
- **CPR Time (post shock or after no shock advised)** – User configurable – 15, 30, 45, 60, 90, 120, 180 seconds
- **Device Date/Time**
- **Voice Prompt Volume** – Allows user to change speaker volume
- **ECG Display (if option purchased)** – Turns display on/off for AED mode

- **Motion Detection** – User defined On/Off (default Off)
- **Service Alert** – Audio alarm if the device needs servicing. Configurable on/off.
- **Manual Access (if option purchased)** – Devices configured with an ECG display may be set up to allow user to initiate a charge and shock without analysis

cprMAX Technology Settings:

- **Initial CPR** – User defined time for CPR after first analysis. Can be set to OFF, 15, 30, 45, 60, 90, 120 and 180 seconds.
- **Preshock CPR** – Allows for CPR while device is charging.
- **Confirmation Analysis** – Confirms shockable rhythm after completion of Initial CPR or Preshock CPR periods and prior to Push to Shock prompt.
- **Stacked Shocks** – (ON/OFF) allows for provision of CPR after each shock
- **Pulse Check** – (ON/OFF) eliminates Pulse Check prompt after shock delivery.

DISPLAY

Backlit LCD displays number of shocks delivered, elapsed time, text and graphics of heart rhythm and optional ECG

Size: 120 mm (4.7 in) x 89 mm (3.5 in)

Frequency Response: 0.55 Hz to 21 Hz (-3 dB), nominal

ECG option:

- **Waveform Sweep Speed** – 25 mm/sec for ECG, nominal
- **Waveform Viewing Time** – Minimum 4 seconds
- **Waveform Amplitude** – 1 cm/mV, nominal
- **Heart Rate** – 20 to 300 BPM digital display, Display "----" if heart rate is less than 20 bpm, heart symbol flashes for each QRS detection

ECG information is received from the adult and Infant/Child pads in anterior-lateral or anterior-posterior positions. A 3-wire cable can be used for ECG monitoring (Lead II).

ENVIRONMENTAL

One Hour Operating Temperature (from room temperature to temperature extreme, one hour duration): -20 to 60°C (-4 to +140°F)

Operating Temperature: 0 to 50°C (32 to 122°F)

Storage Temperature: -30 to 60°C (-22 to +144°F) with battery and electrodes (maximum exposure limited to 7 days)

Atmospheric Pressure: 575hPa to 1060hPa (15,000 to -1250 feet)

Relative Humidity: 5 to 95% (non-condensing)

Dust/Water Resistance: IP55 with battery and REDI-PAK™ electrodes installed (IEC 60529/EN 60529)

Bump: 15 Grms., 1000 bumps (IEC 600-68-2-29)

Shock: 40 G peak, 15 - 23 msec, 45 Hz cross over frequency

Drop: 1 meter drop on each corner, edge and surface (MIL-STD-810F, 516.5, Procedure IV)

Vibration: Random vibration test - MIL-STD-810F, Method 514.5, Category 20; Ground vehicle 3.15 G-rms. 1 hour per axis

EMI:

- **Radiated** - IEC 60601-2-4 EN/IEC60601-1-2, CISPR 11 Class B Group 1
- **Immunity** - EN/IEC60601-2-4, EN/IEC60601-1-2, IEC61000-4-3, Level 3/ RTCA/DO 160D, Sec 20.5, Cat 7/ EN/IEC60601-1-2, IEC61000-4-8, Level 2
- **Aircraft (radiated)** - RTCA/DO 160D, Sec 21.4, Cat M

EVENT DOCUMENTATION AND COMMUNICATION

Memory Capacity: Dual patient storage. Minimum 40 minutes ECG for current patient. Summarized data for previous patient.

Report Types: Continuous ECG, summary (critical resuscitation events and associated ECG waveforms), event log report (report of time stamped entries reflecting operator and device activity), test log report (self test activity report)

Capacity: Minimum 100 time stamped event log entries

Data Review: CODE-STAT™ Suite 6.0 system, DATA TRANSFER™ Express 2.0 system

Communications: Infrared wireless transfer to personal computer

BATTERY AND READINESS DISPLAY

Note: See operating instructions for information on battery care

Primary Battery (nonrechargeable battery with status indicator):

- **Type** – Lithium Manganese Dioxide (Li/MnO₂), 12.0V, 6.2 amp-hours
- **Capacity** – Typically will provide 440 200 joule shocks or 1030 minutes of operating time with a new battery (380 200 joule shocks or 900 minutes of operating time minimum)
- **Weight** – 0.45 Kg (1.0 lb)
- **Shelf Life** – After the battery is stored for 5 years at 20°C to 30°C, the device will provide 48 months of standby life
- **Standby life (assuming daily tests only)** – A new battery provides device power for 5 years
- **Low battery indication** – At least 30 shocks still remain when low battery is indicated

PHYSICAL CHARACTERISTICS

Height: 8.7 cm (3.4 in)

Width: 23.4 cm (9.2 in)

Depth: 27.7 cm (10.9 in)

Weight: XX kg (XX lbs) with one set of REDI-PAK electrodes and one nonrechargeable battery

For further information please call Medtronic at 1.800.442.1142 or visit www.medtronic-ers.com

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