### THE ART AND SCIENCE OF DEFIBRILLATION™



### Lifeline AUTO<sup>®</sup> Fully-Automatic Defibrillator

Defibtech Lifeline AEDs offer industry-leading innovation, simplicity, and elegance. The Lifeline AUTO is a fully-automated defibrillator designed to analyze heart rhythm and automatically deliver a shock - without intervention - if needed.

Technologically advanced enough to include all mission critical features necessary to provide the most advanced treatment for Sudden Cardiac Arrest. Yet so simple and unintimidating to use that virtually anyone can quickly and effectively help save a life.

The Lifeline AUTO was developed by experienced multidisciplinary engineering teams and incorporates state-of-the-art digital signal processing techniques and advanced ECG analysis algorithms. The device exceeds the American Heart Association performance recommendations - giving the user confidence the correct therapy is being delivered.

In the event of a cardiac arrest, defibrillation using an AED within three minutes can increase survival rates from less than 8 percent to more than 70 percent. The Lifeline AUTO defibrillator, with its one-button operation, uses advanced biphasic technology and automatically adjusts the shock delivery to the person's individual needs.

Defibtech AED deployments include workplaces, government buildings, airports and aircraft, rail stations and trains, educational institutions, malls, factories, emergency vehicles, healthcare facilities, resorts, arenas, and marine vessels. More than a quarter-million units have been shipped worldwide.



# defibtech Lifeline AEDs

# Offering the Best Selection for Saving a Life

Defibtech is a leader and innovator in the design and manufacture of automated external defibrillators (AEDs), mechanical chest compressors, and other life-saving resuscitation products. By using advanced design and manufacturing techniques, Defibtech provides value-oriented, easy-to-use solutions with high quality and reliability.

### Life-Saving Design

Defibtech's technologically advanced devices include the Lifeline family of fully-featured AEDs with distinctive yellow hourglass shapes, roomy handles, and rubberized surfaces. Sophisticated enough to meet the needs of the most demanding first responders, they are also incredibly easy for the untrained to use. Virtually anyone can be a lifesaver with a Lifeline AED as it leads the user through a rescue step-by-step.

The Lifeline AED product line includes a semi-automatic defibrillator, a fully-automated defibrillator that analyzes heart rhythms and automatically delivers a shock, an AED capable of an ECG waveform display at the touch of a button, and the first AED with full-motion color video.

Built to exacting medical standards as well as to U.S. Military specifications, Defibtech's Lifeline AEDs are lightweight, robust, dust protected, spray and water resistant, and meet "shock and drop" specifications for use in tough environments. They are also easily maintained and field upgradable, on-site, when CPR guidelines change.

### A Trusted Industry Leader

Defibtech has drawn accolades and won numerous awards for its record of innovative sleek product designs, revenue growth, and commitment to quality and service excellence. Deployments include workplaces, government buildings, airports and aircraft, rail stations and trains, educational institutions, emergency vehicles, resorts, arenas, and waterway vessels.

Headquartered in Guilford, Connecticut, all life-saving products are conceived and developed in-house, and built in the United States in state-of-the-art facilities. For more information about Defibtech and its products, visit www.defibtech.com.

## **Defibtech Lifeline AUTO Fully-Automatic Defibrillator**

TECHNICAL SPECIFICATIONS<sup>†</sup>

#### DEFIBRILLATOR

#### TYPE

Fully-automatic external defibrillator

MODEL DDU-120A, DDU-120E

#### WAVEFORM **Biphasic Truncated Exponential** (Impedance compensated)

**ENERGY** Adult: 150 Joules Child / Infant: 50 Joules (Nominal into 50 Ohm load)

**CHARGE TIME\*** 4 seconds or less (from shock advised)

#### **VOICE PROMPTS**

Extensive voice prompts guide user through operation of the unit

#### **PATIENT ANALYSIS SYSTEM**

#### PATIENT ANALYSIS

Automatically evaluates patient impedance for proper pad contact. Monitors signal quality and analyzes patient ECG for shockable/ non-shockable rhythms.

#### **BATTERY PACK**

СР	R	PA	CII	١G

Metronome CONTROLS

Lighted On/Off button

#### **INDICATORS**

- · "check pads"
- · "do not touch patient" · "analyzing"
- armed-for-shock
- **AED Status LED**

#### **RESCUE PROTOCOL**

AHA/ERC (default); supports protocol updates

\*Typical, with new battery at 25°C

SENSITIVITY/SPECIFICITY

meets AAMI DF80 requirements

IEC-60601-2-4 requirements;

Meets or exceeds

#### **DEFIBRILLATION / MONITORING PADS**

#### MODEL

Adult: DDP-100 Child / Infant: DDP-200P

#### TYPE

Pre-connected, single-use, non-polarized, disposable, self-adhesive electrodes with cable and connector

### **EVENT DOCUMENTATION**

#### **INTERNAL EVENT RECORD**

Critical ECG segments and rescue event parameters are recorded and can be downloaded to a removable data card

#### PC-BASED EVENT REVIEW

ECG with event tag display, and audio playback when available

#### **ENVIRONMENTAL**

#### **TEMPERATURE**

Operating: 0 to 50°C (32 to 122°F) One Hour Operating Temperature Limit (extreme cold): -20°C (-4°F)\*\*\*

#### Standby: 0 to 50°C (32 to 122°F)

**RELATIVE HUMIDITY** Operating / Standby: 5%-95% (non-condensing)

#### ALTITUDE -500 to 15,000 ft (-150 to 4500 m) per MIL-STD-810F 500.4 Procedure II

VIBRATION Ground (MIL-STD-810F 514.5 Category 20) Helicopter (RTCA/DO-160D, Section 8.8.2, Cat R. Zone 2, Curve G) Jet Aircraft (RTCA/DO-160D Section 8. Cat H. Zone 2. Curves B & R)

### **PHYSICAL**

SIZE 8.5 x 11.8 x 2.7 inches (22 x 30 x 7 cm)

WEIGHT (Approximate) With DBP-1400: 4.2 lbs (1.9 kg) With DBP-2800: 4.4 lbs (2.0 kg)

**AUTOMATIC** Automatic daily, weekly, monthly and quarterly circuitry tests

**BATTERY INSERTION** System integrity test on battery insertion

PAD PRESENCE Pads preconnected tested daily

> DAC-A705-EN-BB Issued: 2018-10-03

**USER-INITIATED** Unit and battery pack system test initiated by the user

**STATUS INDICATION** Visual and audible indication of unit status

S

Audible

\*Typical, with new battery at 25°C

#### and event data storage (no audio option) or up to 1 hour and 40 minutes of audio (audio option). ECG and event storage on a re-

**REMOVABLE STORAGE** 

(optional) Up to 12 hours of ECG

SURFACE AREA\*\*

Adult: 16 inches<sup>2</sup> (103 cm<sup>2</sup>)

Child / Infant: 7.75 inches<sup>2</sup> (50 cm<sup>2</sup>)

\*\*Nominal, each pad

movable data card. Actual length of storage is dependent on card capacity.

#### SHOCK / DROP ABUSE TOLERANCE

MIL-STD-810F 516.5 Procedure IV (1 meter, any edge, corner, or surface, in standby mode)

#### **SEALING / WATER** RESISTANCE

IEC 60529 class IP54; Dust Protected, Splash Proof (battery pack installed)

**ESD** EN 61000-4-2 (15kV or direct contact up to 8kV)

**EMC** (Emission) EN 55011 Class B Group 1 and FCC Part 15

**EMC** (Immunity) EN 61000-4-3 (20V/m)

> \*\*\*From room temperature to temperature extreme. one hour duration



MODEL DBP-2800	MODEL DBP-1400		
<b>POWER</b> 15VDC, 2800 mAh	<b>POWER</b> 15VDC, 1400 mAh		
<b>CAPACITY</b> 300 shocks or 16 hours continuous operation*	<b>CAPACITY</b> 125 shocks or 8 hours continuous operation*		
<b>STANDBY LIFE</b> 7 years (installed in AED with 9V ASI battery)*	<b>STANDBY LIFE</b> 5 years (installed in AED with 9V ASI battery)*		
TYPE Lithium/Manganese Dioxide	LOW BATTERY INDICATORS		

Disposable, recyclable, non-rechargeable

**SELF TESTS** 

#### **BRIEF SUMMARY OF INDICATIONS, CONTRAINDICATIONS AND OTHER IMPORTANT SAFETY INFORMATION**

### When should the Defibtech Automated External Defibrillator (AED) be used - what are its indications?

Lifeline/ReviveR DDU-100 and Lifeline/ReviveR AUTO DDU-120 Automated External Defibrillators (AEDs) are indicated for use on victims of sudden cardiac arrest (SCA) who are:

- Unconscious and unresponsive
- Not breathing or not breathing normally

Lifeline/ReviveR DDU-100 and Lifeline/ReviveR AUTO DDU-120 AEDs may be used with Defibtech adult defibrillation pads (model number DDP-100). For patients under 8 years old, or weighing less than 55 lbs (25 kg), use Defibtech child/infant defibrillation pads (model number DDP-200P), if available.

### When should the Defibtech AED not be used - what are its contraindications?

Lifeline/ReviveR DDU-100 and Lifeline/ReviveR AUTO DDU-120 Automated External Defibrillators (AEDs) should not be used if the victim is responsive or conscious.

#### What other information is important about using the AED?

Do not delay therapy to determine exact age or weight. If pediatric pads are not available, apply adult pads in the position as shown for a child/ infant and use the AED.

#### What are the potential adverse health effects of using an AED?

The potential adverse effects (e.g., complications) associated with use of an automated external defibrillator include, but are not limited to, the following:

- Failure to identify shockable arrhythmia.
- Failure to deliver a defibrillation shock in the presence of VF or pulseless VT, which may result in death or permanent injury.
- Inappropriate energy, which could cause failed defibrillation or postshock dysfunction.
- Myocardial damage.
- Fire hazard in the presence of high oxygen concentration or flammable anesthetic agents.
- Incorrectly shocking a pulse sustaining rhythm and inducing VF or cardiac arrest.
- Bystander shock from patient contact during defibrillation shock.
- Interaction with pacemakers.
- Skin burns around the defibrillation pads placement area.
- Allergic dermatitis due to sensitivity to the materials used in the defibrillation pads construction.
- Minor skin rash.

#### What are some of the relevant warnings related to the AED?

- Hazardous electrical output. This equipment is for use only by qualified personnel.
- Possible fire or explosion. Do not use in the presence of flammable gases or anesthetics. Use care when operating this device close to oxygen sources (such as bag-valve-mask devices or ventilator tubing). Turn off gas source or move source away from patient during defibrillation, if necessary.
- The DDU-100 Series AED has not been evaluated or approved for use in hazardous locations as defined in the National Electric Code standard. In compliance with IEC classification, the DDU-100 Series AED is not to be used in the presence of flammable substance/air mixtures.
- Improper maintenance can cause the DDU-100 Series AED not to function. Maintain the DDU-100 Series AED only as described in the User Manual and Operating Guide. The AED contains no userserviceable parts — do not take the unit apart.
- Do not open sealed pads package until pads are to be used. The packaging should be opened only immediately prior to use, otherwise the pads may dry out and become non-functional.
- Do not touch the patient during defibrillation. Defibrillation current can cause operator or bystander injury.
- The defibrillation pads are intended for one-time use only and must be discarded after use. Reuse can lead to potential cross infection, improper performance of the device, inadequate delivery of therapy, and/or injury to the patient or operator.
- CPR during analysis can cause incorrect or delayed diagnosis by the patient analysis system.
- User-initiated and automatic self-tests are designed to assess the DDU-100 Series AED's readiness for use. However, no degree of testing can assure performance or detect abuse, damage, or a defect that occurred after the most recent test is completed.
- Even if defibrillation occurs, the sudden cardiac arrest event may not result in survival.

#### What are some of the relevant cautions related to the AED?

- Follow all battery pack labeling instructions. Do not install battery packs after the expiration date.
- Follow all defibrillation pad label instructions. Use defibrillation pads prior to their expiration date.
- Use and store the DDU-100 Series AED only within the range of environmental conditions specified in the technical specifications.

 $\ensuremath{\textit{Caution:}}$  Federal (U.S.A) law restricts this device to sale by or on the order of a physician.

Please refer to the Operating Guide provided with your AED for user instructions, complete list of warnings and cautions, operator training requirements, summary of primary clinical studies, technical specifications, and other important information. The Operating Guide, for concise guidance on set-up, use, maintenance and technical specifications, and User Manual, for comprehensive training on set-up, use and maintenance; and source for complete technical specifications, are also available at www.defibtech.com/support.



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