

STATE OF NEW HAMPSHIRE
Dept. of Administrative Services
Div. of Procurement and Support Services
Bureau of Purchase and Property
State House Annex
Concord, New Hampshire 03301

Date: July 6, 2020

NOTICE OF CONTRACT

COMMODITY: Automatic External Defibrillators (AED's)
Defibtech

CONTRACT NO.: 8002699

NIGP: 465-1400

VENDOR: LifeSavers Inc **VENDOR # :** VC #166730
39 Plymouth Street
Fairfield, NJ 07004

CONTACT PERSON: Bob Stickel
Tel. No.: 973-244-9111 or 866-641-1200
Fax No.: 973-244-1666
E-Mail: Bob@lifesaversinc.com

EFFECTIVE FROM: July 1, 2020 **Through:** June 30, 2023

Prior to placing an order, please contact Bill Wood, NH Department of Safety, Bureau of Emergency Medical Services, at (603)-223-4228 or William.Wood@dos.nh.gov for guidance in the products being ordered.

AED's in New Hampshire must be registered with the NH Bureau of Emergency Medical Services.

DELIVERY TIME: Ten (10) Business days from placement of order.

F.O.B.: Destination **TERMS:** Net 30

ORDERING: By telephone or fax (see above)

MINIMUM ORDERS: There is no minimum order required under this contract.

INVOICING & PAYMENTS: All invoicing is to reflect contract number and all invoices are to be sent to the requesting agency.

Warranty: For a period of not less than the manufacturer's standard period of time, or five (5) years, whichever is longer, from the date the items are received, inspected and accepted by the State of New Hampshire.

QUESTIONS: Loretta Razin at Loretta.Razin@DAS.NH.GOV or call 603-271-0579

DELIVERY LOCATIONS:

Current State of New Hampshire agency/institution locations which, if you are awarded a contract, you are expected to service.

SPECIFICATION COMPLIANCE:

Unless otherwise specified by the Bureau of Purchase and Property, all equipment offered by the Contractor must be new; shall not be used, rebuilt, refurbished; shall not have been used as demonstration equipment, and shall not have been placed anywhere for evaluation purposes.

Automated External Defibrillators (AED's) shall be in accordance with the following specifications:

- Shall meet the 2010 American Heart Association/Emergency Care Committee (AHA/ECC) cardiopulmonary resuscitation requirements. The AED's shall be capable of being upgraded to future AHA/ECC requirements by the user.
- US Food and Drug Administration (USFDA) approved
- Bi-Phasic Shock Delivery
- AED's shall be capable of having cardiac arrest events data downloaded for post-event review. (the actual download equipment with contract pricing shall be optional for purchase by the AED agency)
- Prompts that is visual and auditory (in English) providing the user with AED uses instructions. These prompts must provide detailed directions with one or more pictures /diagrams showing user best/most appropriate locations for defibrillation pad(s) placement.
- Elapsed Time
- Internal capability to record and abstract with a minimum of 15 minutes recording time of an incident event
- Pre-approved "Medical Authorization" form for USFDA compliance (if necessary) for ordering organization contact information
- Printed Operator's Guide (in English)
- 1 (one) set Adult Defibrillation Pad(s) with a minimum of a 2-year expiration date. Defibrillation Pad(s) shall be pre-gelled, self-adhesive, one-time use and disposable
- Lithium battery with an operational user life of a minimum of four years
- Pediatric patient capability
- Device shall be able to perform: daily automatic self-tests (to include, but not limited to: internal test, circuitry, waveform delivery system, defibrillation pad(s) connection status and battery status
 - Battery Insertion Test – upon battery insertion, automatic internal self-test for device readiness with visual or auditory status update.
 - Green "ready" status indicator identifying device available for use. Audible and/or visual indicator identifying need for device maintenance
- 24 hour per day Vendor or Manufacturer technical support