22-1365-01

An act relating to liability; creating s. 768.1325, F.S.; providing immunity from civil liability for certain persons acquiring or using automatic external defibrillator devices; repealing s. 768.13(4), F.S., relating to automatic external defibrillators; amending s. 401.2915, F.S.; deleting a requirement that persons having access to automatic external defibrillators obtain specified training; providing an effective date.

A bill to be entitled

WHEREAS, over 700 lives are lost every day to sudden cardiac arrest in the United States alone, and

WHEREAS, two out of every three sudden cardiac deaths occur before a victim can reach a hospital, and

WHEREAS, more than 95 percent of these cardiac arrest victims will die, many because of the lack of readily available lifesaving medical equipment, and

WHEREAS, with current medical technology, up to 30 percent of cardiac arrest victims could be saved if victims had access to immediate medical response, including defibrillation and cardiopulmonary resuscitation, and

WHEREAS, once a victim has suffered a cardiac arrest, every minute that passes before returning the heart to a normal rhythm decreases the chance of survival by 10 percent, and

WHEREAS, most cardiac arrests are caused by abnormal heart rhythms called ventricular fibrillation, which occurs when the heart's electrical system malfunctions, causing a

chaotic rhythm that prevents the heart from pumping oxygen to the victim's brain and body, and

WHEREAS, communities that have implemented programs ensuring a widespread public access to defibrillators, combined with appropriate training, maintenance, and coordination with local emergency medical systems, have dramatically improved the survival rates from cardiac arrest, and

WHEREAS, automatic external defibrillator devices have been demonstrated to be safe and effective, even when used by lay people, since the devices are designed not to allow a user to administer a shock until after the device has analyzed a victim's heart rhythm and determined that an electric shock is required, and

WHEREAS, increased public awareness regarding automatic external defibrillator devices will greatly facilitate their adoption, and

WHEREAS, limiting the liability of users and acquirers of automatic external defibrillator devices in emergency situations may encourage the use of the devices and result in saved lives, NOW, THEREFORE,

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Be It Enacted by the Legislature of the State of Florida:

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Section 1. Section 768.1325, Florida Statutes, is created to read:

28 civil liability.--

liability.-(1) DEFINITIONS.--As used in this section, the term:

768.1325 Cardiac Arrest Survival Act; immunity from

30 <u>(a) "Perceived medical emergency" means circumstances</u>
31 in which the behavior of an individual leads a reasonable

person to believe that the individual is experiencing a life-threatening medical condition that requires an immediate medical response regarding the heart or other cardiopulmonary functioning of the individual.

- (b) "Automatic external defibrillator device" means a
  defibrillator device that:
- 1. Is commercially distributed in accordance with the Federal Food, Drug, and Cosmetic Act;
- 2. Is capable of recognizing the presence or absence of ventricular fibrillation and of determining without intervention by the use of the device whether defibrillation should be performed;
- 3. Upon determining that defibrillation should be performed, is able to deliver an electrical shock to an individual; and
- 4. In the case of a defibrillator device that may be operated in either an automatic or a manual mode, is set to operate in the automatic mode.
- (c) "Harm" means damage or loss of any and all types, including, but not limited to, physical, nonphysical, economic, noneconomic, actual, compensatory, consequential, incidental, and punitive damages or losses.
- (2) IMMUNITY.--Notwithstanding any other provision of law to the contrary, and except as provided in subsection (3), any person who uses or attempts to use an automatic external defibrillator device on a victim of a perceived medical emergency is immune from civil liability for any harm resulting from the use or attempted use of such device, or any act or failure to act in providing or arranging further medical treatment; in addition, any person who acquired the

device is immune from such liability, if the harm was not due to the failure of such acquirer of the device:

- (a) To notify local emergency response personnel or other appropriate entities of the most recent placement of the device within a reasonable period of time after the device was placed;
  - (b) To properly maintain and test the device; or
- (c) To provide appropriate training in the use of the device to an employee or agent of the acquirer when the employee or agent was the person who used the device on the victim, except that such requirement of training does not apply if:
- 1. The employee or agent was not an employee or agent who would have been reasonably expected to use the device; or
- 2. The period of time elapsing between the engagement of the person as an employee or agent and the occurrence of the harm (or between the acquisition of the device and the occurrence of the harm in any case in which the device was acquired after such engagement of the person) was not a reasonably sufficient period in which to provide the training.
- (3) INAPPLICABILITY OF IMMUNITY.--Immunity under subsection (2) does not apply to a person if:
- (a) The harm involved was caused by that person's willful or criminal misconduct, gross negligence, or reckless misconduct or a conscious, flagrant indifference to the rights or safety of the victim who was harmed;
- (b) The person is a hospital, clinic, or other entity whose primary purpose is providing health care directly to patients and the harm was caused by an employee or agent of the entity who used the device while acting within the scope of the employment or agency of the employee or agent; or

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- (c) The person is an acquirer of the device who leased the device to a health care entity (or who otherwise provided the device to such entity for compensation without selling the device to the entity) and the harm was caused by an employee or agent of the entity who used the device while acting within the scope of the employment or agency of the employee or agent.
- (4) CAUSE OF ACTION, DUTY NOT ESTABLISHED. -- This section does not establish any cause of action. This section does not require that an automatic external defibrillator be placed at any building or other location or require an acquirer to make available on its premises one or more employees or agents trained in the use of the device.
- Section 2. Subsection (4) of section 768.13, Florida Statutes, is repealed.
- Section 3. Section 401.2915, Florida Statutes, is amended to read:
- 401.2915 Automatic external defibrillators.--It is the intent of the Legislature that an automatic external defibrillator may be used by any person for the purpose of saving the life of another person in cardiac arrest. In order to ensure public health and safety:
- (1) All persons who have access to or use an automatic external defibrillator must obtain appropriate training, to include completion of a course in cardiopulmonary resuscitation or successful completion of a basic first aid course that includes cardiopulmonary resuscitation training, and demonstrated proficiency in the use of an automatic external defibrillator;
- (2) Any person or entity in possession of an automatic 31 external defibrillator is encouraged to register with the

local emergency medical services medical director the existence and location of the automatic external defibrillator; and (3) Any person who uses an automatic external defibrillator is required to activate the emergency medical services system as soon as possible upon use of the automatic external defibrillator. Section 4. This act shall take effect October 1, 2001. SENATE SUMMARY Provides immunity from civil liability for persons who use or attempt to use automatic external defibrillator devices in perceived medical emergencies, as defined. Also provides immunity to the acquirers of such devices under specified circumstances. However, immunity does not attach in cases of willful or criminal action, gross negligence, or indifference to the rights or safety of the person harmed; if the person is a hospital, clinic, or similar entity; or if the person is an acquirer of a device who leased it to a health care entity.