SOMA TECH INTL

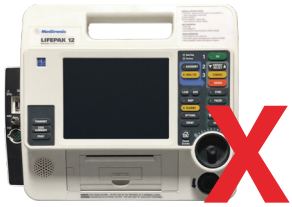
ARE YOUR DEFIBRILLATORS FDA APPROVED?

The FDA requires AED and defibrillator manufacturers to obtain premarket approval (PMA) for all models they sell. To obtain a PMA, the manufacturer must prove that its devices are safe, effective, and reliable. If a device, or its accessories, has not received a PMA it will no longer be available from the manufacturer after **February 3, 2022**. If your facility is still utilizing unapproved devices, now is the time to explore alternative options to ensure that the replacement AEDS and defibrillators are in place by the deadline.

IF YOU HAVE THESE IT'S TIME TO UPGRADE!

LIFEPAK 20

ZOLL M



LIFEPAK 12

UPGRADE TO THESE BEFORE FEBRUARY 3, 2022

LIFEPAK 20E

ZOLL R

LIFEPAK 15

SAME SERVICE & WARRANTY **AS NEW**

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UP TO 50% BELOW OEM

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