

Dana Ingram presents Defibrillator and AED Best Practices: Saving lives, one test a time

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Wednesday, March 11, 2015, 2:00pm EST

PRESENTER: DANA INGRAM, PRODUCT MANAGER FOR FLUKE BIOMEDICAL



As one of Fluke Biomedical's Product Managers, Dana Ingram manages the Defibrillator, AED and Pacemaker analyzers and testers product line. Dana has over 7 years of product marketing experience, as she was previously a Product Manager for the Global Strategy & Development Infusion Device Group at CareFusion in San Diego. Prior to that, she held several Product Marketing positions within Mentor Worldwide, a Johnson & Johnson Company.

In this 60 minute presentation Dana will be discussing the best ways to test defibrillators and AEDs for preventive maintenance. This will include information on current global regulatory standards.

With high AED failure rates, there has been increased FDA scrutiny in testing defibrillators and AEDs. Join Fluke Biomedical's Product Marketing Manager, Dana Ingram, as she discusses how to ensure proper performance of modern defibrillators and AEDs, including:

- · Global testing standards
- Best practices in defibrillators
- Best practices in AEDs

Get valuable insights on one of the most highly regulated medical devices and learn how we can save lives by making sure defibrillators and AEDs work the first time, every time.

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Defibrillator and AED Webinar Workbook (Fill-in-the-Blank)

1)	Defibrillators and AEDs convert certain	and	back to normal	
2)	Regulated by the FDA, defibrillators are classified	d as a	medical device	
3)	The standards that require defibrillators to be tested on different resistance loads to ensure proper			
	current is delivered to patients with different imp	pedances are	and	
4)	Theestablished	standards for particular b	asic safety and essential	
	performance requirements for cardiac defibrillat	ors.		
5)	Preventative maintenance testing on defibrillator	rs is recommended	times a year	
6)	The standard testing load for defibrillator devices	s is approximately	ohms.	
7)	Check and test performance of all patient parame			
8)	Human impedance variability has been shown to			
9)	Recall/corrective action root causes are		, and	
10) Many AED failures could have been solved by pe	rforming		
	and	·		
11	are under increased s	crutiny by the USA FDA dı	ue to high failure incident	
	reporting rates.			

ANSWERS: 1.) heart arrhythmias and ventricular fibrillation, 2.) Class III, 3.) IEC 60601-2-4 and AAMI DF80, 4.) IEC (International Electrotechnical Commission), 5.) Two (2), 6.) 50, 7.) ECG, Arrhythmia, Sp02, NIBP, IBP, ETCO2, 8.) 25 ohms to 180 ohms, 9.) microchip failure, battery failure, and power supply/recharging circuit failure, 10.) daily checks, monthly tests, and detailed tests, 11.) AEDs

Thank you for attending Webinar Wednesday!

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