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PHILIPS EMERGIN RECEIVES FDA 510(K) CLEARANCE TO MARKET INTELLISPACE EVENT MANAGEMENT SOLUTION

Philips Emergin received 510(k) clearance from the U.S. Food and Drug Administration on January 4, 2011 for the Philips Emergin IntelliSpace Event Management solution. This solution, available in Spring 2011, builds on the success of the Emergin platform to provide supplemental notifications and escalations of clinical alarms and alerts to patient care providers through their preferred communication device.

Philips Emergin has gained wide acceptance among caregivers and hospital chief information officers (CIOs) for its innovation and ongoing commitment to excellence. This latest achievement reassures hospitals that Philips Emergin will continue to address the rigorous standards necessary to function as a trusted partner in the healthcare industry.

The Philips Emergin IntelliSpace Event Management solution will provide an interoperable platform that integrates with a broad suite of medical devices, enterprise nurse call systems and output communication devices to standardize management of hospital alarm and alert notifications. It dispatches supplemental notifications from medical devices such as patient monitors, bed systems, and infusion pumps to a variety of output communication devices such as one-way pagers, wireless smartphone devices, and other mobile communication devices.

For more information, visit Philips booth #3845 during the HIMSS Annual Conference in Orlando February 21-23.