News Release

Contact: Larry LeBlanc
Phone: +1 717-309-7006
lleblanc@deviceevents.com

For Immediate Release

Device Events selected to participate in HealthTrust 2017 Innovations Summit

York, PA USA — 12 July 2017 – Device Events announced today that it had been selected to participate in the 2017 **HealthTrust Innovation Summit** on October 5-6 in Ponta Vedra Beach, Florida.

HealthTrust selects companies from many applicants to participate in the Innovation Summit to present "truly innovative products to service line clinical experts".

Founded in 1999, HealthTrust members include 1,600 hospitals and more than 26,000 non-acute care sites in the U.S. and U.K. It offers the industry's only national committed model GPO.

Madris Tomes
Founder & CEO
Device Events, LLC

Device Events Inc., founded by Ms. Madris Tomes in 2015, provides a cloud-based software service that enables healthcare organizations to detect and analyze problems with medical devices early – before FDA recalls. The service provides powerful search, retrieve and reporting on the FDA MAUDE database, email notifications of adverse event activity and trend visualizations to help users make informed decisions on purchases and continued use of potentially problem devices

"There are over 6 million adverse event reports in the MAUDE database and more than 65,000 new reports are added every month. It can take the FDA, 2 months to 2 years to identify a problem device and act on it. During this time, the healthcare community is usually unaware and continue to use these devices," comments Tomes, Founder and CEO of Device Events.

"Over the past 18 months, we've developed a very powerful decision support tool for healthcare organizations to make better and earlier decisions on problem medical devices that can critically impacts safety and health outcomes for patients and the financial risk for organizations," notes Ms. Tomes.

About Device Events Inc.
Device Events proactively identifies patterns of problems with medical devices.
Healthcare organizations, insurance companies, manufacturers, and patient advocacy groups alike can utilize Device Events to quickly identify problematic devices and understand the history, quantity, and severity of related adverse events. With the service, users make more informed decisions, resulting in better health outcomes and reduced risk for both patients and their organizations. To learn more visit www.deviceevents.com.

####