

Weitz and Luxenberg Investigating Injuries Caused by Recalled Beacon Tip Angiographic Catheters from Cook Medical

NEW YORK, N.Y., Aug. 12, 2015 (SEND2PRESS NEWSWIRE) – Weitz & Luxenberg, P.C., today launched an investigation into injuries sustained by patients when catheters used during cardiac angiogram procedures to inject contrast dye into blood vessels malfunctioned by either splitting or separating, the firm announced.

The probe centers around one particular brand of angiographic catheters made by Cook Medical of Bloomington, Indiana, Weitz & Luxenberg said. The recall, classified as a Class I recall by the FDA, notes these problems could “cause serious injury to the patient and require additional medical intervention to retrieve the tip, or cause death.”

A cardiac angiogram is an imaging procedure that allows doctors to measure and visualize blood flow within the heart. The American Heart Association (AHA) describes the procedure as typically having minimal side effects, Weitz & Luxenberg said.

The Cook Medical Beacon Tip angiographic catheters at the center of the investigation have been reported to either split or break at the tip while inside the patient’s vasculature, the firm said.

Weitz & Luxenberg began its investigation after Cook Medical on July 2 recalled 38,895 domestically distributed units of its Beacon Tip Angiographic Catheters. The recalled catheters were distributed between June 2013 and June 2015, Weitz & Luxenberg said.

The recalled products include the Beacon Tip Torcon NB Advantage Catheters (Cook Medical catalog prefix HNBR5.0), Beacon Tip Royal Flush Plus High-Flow Catheters (catalog prefix HNR4.0) and Slip-Cath Beacon Tip Catheters (SCBR5.0), Weitz & Luxenberg said.

The voluntary recall was posted by the U.S. Food and Drug Administration (FDA) on August 7. The FDA explained the recall was precipitated by Cook Medical receiving 26 reports of catheter malfunctions, 14 of which involved adverse events.

“The consequences of a catheter malfunctioning inside a heart chamber or a blood vessel during an angiogram could be potentially devastating,” said Paul Pennock, managing attorney of Weitz & Luxenberg’s Defective Drugs and Devices unit.

“As defined by the FDA, a serious adverse event may mean that use of the

medical product substantially disrupted the patient's ability to conduct normal life functions, created a substantial risk of death at the time of the adverse event, or perhaps even resulted in death," he added.

Exploring Injured Heart Patients' Legal Rights:

Weitz & Luxenberg said its investigation is for now focused on meeting with patients who were injured by Cook Medical Beacon Tip angiographic catheters.

Weitz & Luxenberg said that it also plans to meet with representatives of the estates of such patients who died as a result of being harmed by the Cook Medical Beacon Tip angiographic catheters.

The firm indicated that one purpose of these meetings will be to help the injured patients or their estates understand their legal rights to compensation from Cook Medical.

According to Weitz & Luxenberg, harm caused by a malfunctioning Cook Medical Beacon Tip angiographic catheter may give rise to a right to recover medical care expenses related to post-injury treatment.

Weitz & Luxenberg said the injured patients or their estates may also be entitled to recover lost wages if medical interventions to treat a Cook Beacon Tip angiographic catheter injury caused the patient to be absent from his or her place of employment or to be laid off.

The firm said it is also possible an injured patient or his or her estate may be able to receive compensation for other damages spawned by the malfunction of a Cook Beacon Tip angiographic catheter.

"If you were injured during a cardiac angiogram due to breakage of this device, or if loved ones suffered because of your injury, then you may have legal rights against Cook Medical," said Ellen Relkin, of counsel attorney in Weitz & Luxenberg's Defective Drugs and Devices unit.

To arrange a free consultation, contact Weitz & Luxenberg toll-free at 800-476-6070 or online at <http://www.weitzlux.com/>.

About Weitz & Luxenberg:

Weitz & Luxenberg, P.C. is among the nation's leading and most readily recognized personal injury and consumer protection law firms. Weitz & Luxenberg's numerous litigation areas include: mesothelioma, defective medicine and devices, environmental pollutants, products liability, consumer protection, accidents, personal injury, and medical malpractice. Victims of consumer fraud are invited to rely on Weitz & Luxenberg's more than 25 years of experience handling such cases. You can contact the firm's Client Relations department at 800-476-6070 or at clientrelations@weitzlux.com.

More information: <http://www.weitzlux.com/>.

Twitter: @WeitzLuxenberg

News issued by: Weitz and Luxenberg P.C.



Send2Press® Newswire

Original Image:

<https://www.send2press.com/wire/images/15-0803-weitz-luxenberg-500x375.jpg>

#

Original Story ID: 2015-0812-02 (10423) :: weitz-and-luxenberg-investigating-injuries-caused-by-recalled-beacon-tip-angiographic-catheters-from-cook-medical-2015-0812-02

Original Keywords: Beacon Tip Torcon NB Advantage Catheters, cardiac angiogram procedures Weitz and Luxenberg P.C. New York New York NEW YORK, N.Y.

Alternate Headline: Weitz and Luxenberg Investigating Injuries Caused by Recalled Cook Medical Angiographic Catheters

NEWS ARCHIVE NOTE: this archival news content, issued by the news source via Send2Press Newswire, was originally located in the Send2Press® 2004-2015 2.0 news platform and has been permanently converted/moved (and redirected) into our 3.0 platform. Also note the story "reads" counter (bottom of page) does not include any data prior to Oct. 30, 2016. This press release was originally published/issued: Wed, 12 Aug 2015 13:32:53 +0000

Original Shortcode for Story: <https://i.send2press.com/h6Sf7>