

US Drug Watchdog Now Expands Their Transvaginal Mesh Failure Initiative For All US Women Recipients & They are Offering the Names and Contacts of the Best Women Attorneys

The US Drug Watchdog is expanding their initiative focused on helping women recipients of a failed transvaginal mesh, or a sling because the potential number of US women involved, and because of the severity of the damages. According to the New York Times, in 2010, about 185,000 women underwent procedures in which mesh was implanted vaginally to treat urinary incontinence. That is 2010 alone. Transvaginal mesh failure occurs when the mesh fails to bind to a woman's pelvic tissues. As a consequence of this, the body rejects the mesh that has been sewn into it. As a result, the transvaginal mesh may start to poke its way through the vaginal wall and protrude out of the vaginal tissue. This complication is more commonly referred to as "erosion." Erosion may result in extreme pain, infection, bleeding, vaginal discharge, with no possibility for sexual intercourse. The US Drug Watchdog is urging any woman, who has had the transvaginal mesh, or sling surgical procedure done, and now has developed severe complications to call them anytime at 866-714-6466. As part of their initiative the US Drug Watchdog is offering to send all US women, who are victims of a transvaginal mesh, or sling failure to national caliber attorneys-who are all women. <http://USDrugWatchdog.Com>

([PRWEB](#)) October 02, 2012 -- The US Drug Watchdog is offering to help all US women, who are now victim's of a transvaginal mesh, or sling failure get to the best possible attorneys, and the group is promising the attorneys will all be women. No other Group is offering a service like this. The US Drug Watchdog is saying, "Transvaginal mesh, or sling failures are the most horrific medical device failures we have ever heard of, especially if the failure involves erosion, and there is nothing we will not do to help what we expect to be thousands of US victims." According to court records, transvaginal mesh products were designed for women, who were suffering from pelvic organ prolapse, and or stress urinary incontinence. According to the US FDA the procedure to install a transvaginal was supposed to be minimally-invasive. The US Drug Watchdog now says, "The FDA is now saying transvaginal mesh failures is causing complications in patients such as erosion and infection. Women who experience a transvaginal mesh failure may need to have numerous corrective surgeries, IV therapy, blood transfusions and drainage of hematomas or abscesses. For more information victims of a transvaginal mesh are urged to contact the US Drug Watchdog anytime at 866-714-6466, for the names, and specific contacts of national caliber attorneys-who are all women. <http://USDrugWatchdog.Com>

The US Drug Watchdog is indicating symptoms of a vaginal mesh implant failure may include:

- * Pain during sexual intercourse caused by erosion.
- * Mesh erosion through the vagina (the mesh is actually protruding out of the vaginal wall)
- * Bleeding
- * Vaginal Infection
- * Urinary problems
- * Organ perforation

The US Drug Watchdog says, "One of the biggest problems we have with respect to our failed medical device or recalled drug initiative work is the average US consumer never hears about a drug recall or failures involving medical devices like the Transvaginal Mesh Disaster. If you have a friend, or loved one who is a recipient of a

Transvaginal Mesh that has already failed or is showing symptoms of a failure, please have them call us at 866-714-6466. We want to make certain these Transvaginal Mesh Disaster victims get to the best possible attorneys, to ensure they get the best possible compensation for their ordeal, and we will do our absolute best to make certain all of the attorneys are women." <http://USDrugWatchdog.Com>

U.S. District Court for the Southern District of West Virginia Multidistrict litigation (MDL) consolidates all similar cases into one courtroom and will include the Defendants:

C.R. Bard, Inc., (MDL No. 2187);

American Medical Systems Inc., MDL No. 2325);

Boston Scientific Corp., (MDL No. 2326); and Johnson & Johnson's Ethicon, Inc. (MDL No. 2327)

**Contact Information**

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Online Web 2.0 Version

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