



رقم المحفوظات: ٥٧/٢٥  
رقم الصادر: ١٢/١/٢٠١٣  
بيروت، في: ٣ تشرين الأول ٢٠١٣

جانب نقيب المستشفيات الخاصة في لبنان

**الموضوع:** إشعار بمتابعة جهاز طبي مغروس

Implants, non active, vascular haemostasis device sponge. LeGoo endovascular occlusion gel.

**الجهاز المعنى بالمتابعة:**

- Implants, non active, vascular haemostasis device sponge. LeGoo endovascular occlusion gel.
- Trade Mark: Genzyme Diagnostics Div Genzyme Corp
- Local Representative:

نرجو الاطلاع على التوصية الصادرة عن وكالة

Medicine and Health Care Products Regulatory Agency (UK) MHRA

والتوصية الصادرة عن الشركة المصنعة والتي تشير الى سحب هذا المنتج من التداول، نرجو منكم تعميم هذه النشرة على جميع المستشفيات المعنية.

**مرفق ربطا:**

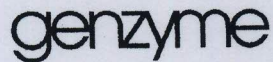
- تقرير MHRA

**يبلغ:**

- دائرة البرامج والمشاريع
- المستشفيات الحكومية
- المحفوظات

مدير عام الصحة  
د. وليد عمار





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Urgent Field Safety Notice  
LeGoo Endovascular Occlusion Gel  
FSCA Reference Number: CR01450  
Type of action – Product Removal

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Date: June 19, 2013

Attention: //////////////////////////////////

**Details on affected devices:**

LeGoo Endovascular Occlusion Gel

Catalog #'s

- 10-0025, 10-0050, 10-0100, 10-0250, 10-0500, 10-1000
- LG50IT, LG100IT, LG250IT

All Lots

**Description of the problem:**

The recent implementation of additional enhanced visual inspection measures for work in-process material resulted in detection of particulate matter, primarily fibers, in filled LeGoo Endovascular Occlusion Gel syringes. The identity of the fibers (up to 3 mm in size) included acrylic, polyester, nylon paper, and cotton. In consideration of the controls and processes that had been in place for prior manufactured lots, it has been determined that Genzyme cannot provide with sufficient assurance that LeGoo material distributed to the market is essentially free of particulate matter.

LeGoo is a water-soluble, biocompatible polymer with reverse thermo-sensitive properties allowing it to be injected as a soft viscoelastic gel that forms an elastic plug within the vessel. LeGoo is indicated for temporary endovascular occlusion of blood vessels up to 4 mm in diameter below the neck.

Based on the review of QA investigation results to date, Adverse Events in Safety Database, Complaints in the complaints database, and Literature references it can be concluded that the theoretical medical risk associated with the potential deviation is remote when exposed to the defect. The theoretical risk of introduction of fiber particles in the bloodstream could include inflammation, foreign body reaction, blockage of blood vessels, thrombosis and injury to tissue and/or organs. Given the potential of syringes to contain particulate matter, a FSCA is being initiated.

