Anapharm Europe's First FDA Inspection

Barcelona, April 27, 2016

Anapharm Europe is proud to announce that it has successfully undergone its first US FDA inspection at its Barcelona based bioanalytical lab.
Anapharm Europe is also registered in accordance with FDA's "Generic Drug User Fee Amendments of 2012" requiring "Self-Identification of Generic Drug Facilities, Sites and Organizations".
This inspections will allow Anapharm to strengthen its position in North America and continue to help its customers achieve their registration goals in the United States.

About Anapharm Europe

Over the past few years Anapharm Europe has established itself as a world class provider of bioanalytical services to international Sponsors with its strategically located, GLP-certified, ANVISA certified, GCP-compliant and FDA inspected laboratory in Barcelona. With a wide experience and a successful regulatory history, Anapharm Europe has become a stable analytical partner for its clients. Anapharm Europe's main strengths include its strong expertise for method development and extensive portfolio of validated analytical methods, its high throughput capacity, as well as its top quality performance to deliver results within rigorous timelines at competitive prices.

For further details about Anapharm Europe, please visit www.anapharmeurope.com or contact your Business Development representative at +34 93 223 8636

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