Lotus Labs, why India's No 1 CRO??

Hi,

Government to frame guidelines for companies to set up pharmacovigilance system

In order to strengthen the pharmacovigilance system for collection, processing and forwarding of Adverse Drug Reaction reports (ADRs), Government has formed a high level committee to prepare guidelines for manufacturers on the same to effectively monitor drug safety.

Source: http://www.pharmabiz.com/NewsDetails.aspx?aid=87099&sid=1

US FDA working to make clinical trials less wasteful

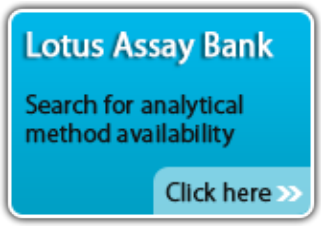
The US FDA is looking to set up master protocols that can continuously run and help cut down on waste that's prevalent in the clinical trial industry, Director of FDA's Center for Drug Evaluation and Research Janet Woodcock said in an interview.

Source: http://www.outsourcing-pharma.com/Clinical-Development/US-FDA-working-to-make-clinical-trials-less-wasteful

Government to ask MCI for sensitizing clinicians, hospitals to collect ADRs for all drugs soon

In order to develop effective reporting of adverse drug reactions (ADRs) in the country, government is planning to sensitize healthcare practitioners in consultation with the Medical Council of India (MCI) as a step towards quantifying ADR data for comprehensive analysis of drug behavior in the Indian population.

Source: http://www.pharmabiz.com/ArticleDetails.aspx?aid=86467&sid=1



About us: Lotus Labs is a Clinical Research Organization which works towards widening the frontiers of Medicine through services of the highest Quality that conform to International Standards, combining speed, accuracy and reliability to be the most admired Clinical Research Organization from India.