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Indian pharma expects US FDA's SUPAC Mfg Eqpt Addendum to boost global regulatory compliance

The Indian pharma industry expects the US FDA's recently released SUPAC (scale-up and post-approval changes): Manufacturing Equipment Addendum will further give fillip to faster clearances in plant audit and product approval. After releasing the Addendum, the US regulatory authority is seeking comments from the pharma industry before June 30, 2013.

Source:

<http://www.pharmabiz.com/NewsDetails.aspx?aid=74732&sid=1>

DCGI instructs CDSCO zonal offices to constitute expert panels to inspect clinical trials

After launching a number of measures in the recent past to closely monitor the clinical trials in the country, the Drugs Controller General of India (DCGI) has asked all the zonal offices of Central Drugs Standard Control Organisation (CDSCO) to set up expert committees to mount regular inspections at the trial sites.

Source:

<http://pharmabiz.com/ArticleDetails.aspx?aid=75142&sid=1>

ICMR to evaluate 800 outstanding traditional practices for herbal drug discovery

With a view to enhancing the rate of herbal drug discovery by tapping into the huge potential in the area traditional practices, the Indian Council for Medical Research (ICMR) is planning to scientifically evaluate at least 800 selected outstanding practices in the next few years. The ICMR has already tied up with the National Innovation Foundation (NIF) which is engaged in the scouting of traditional practices and grassroots innovations. NIF has a database of approximately 80,000 traditional herbal practices from over 540 districts.

Source :

<http://www.pharmabiz.com/NewsDetails.aspx?aid=75303&sid=1>

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