



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

Hi %BASIC:FIRST_NAME%,

DCGI lays down special conditions to allow import of drugs with residual life less than mandated 60%

Only under special conditions like charity activities and national health programmes, the drug regulatory authorities would permit the import of drugs with residual shelf life less than 60 per cent, the office of the Drugs Controller General of India (DCGI) has clarified.

Source: <http://www.pharmabiz.com/NewsDetails.aspx?aid=81655&sid=1>

IPA-US FDA workshop stresses regular upgradation, strengthening of facilities to meet regulatory compliance

The Indian Pharmaceutical Alliance (IPA)-United States Food and Drugs Administration (US FDA) workshop held in Hyderabad stressed on the need for the Industry to continuously upgrade and strengthen manufacturing facilities to face zero tolerance from regulators. According to Dr. Albinus, Deputy Country Director (India), US FDA, the regulators are not biased to any country or company.

Source: <http://www.pharmabiz.com/NewsDetails.aspx?aid=81730&sid=1>

DCGI to explore potential of replacing drug testing on animals with alternate test methods

The Drugs Controller General of India (DCGI) is seriously contemplating to explore the possibilities of replacing drug testing on animals with alternative no animal testing methods. This move aims at curbing the misuse and torture of animals used for animal testing and for safeguarding their interests as ethically required.

Source: <http://www.pharmabiz.com/NewsDetails.aspx?aid=81632&sid=1>

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