

Hi %BASIC:FIRST_NAME%,

US FDA calls industry developing Rheumatoid Arthritis drugs to focus on dosing regimen, efficacy

US FDA is now insisting to the pharma industry which is engaged in the drug development of rheumatoid arthritis (RA) to focus on three aspects. These are dosing regimen, efficacy factors and drug-device combination product considerations.

Source: <http://pharmabiz.com/NewsDetails.aspx?aid=77032&sid=1>

Need for insurance solutions for liability risks in life sciences industry in India: Expert

Spurred with the recent events of a spate of regulatory actions against manufacturers in India, product liability insurance has become relevant, taking into consideration existing risk environment globally coupled with shrinking risk insurance market in India.

Source: <http://pharmabiz.com/PrintArticle.aspx?aid=76714&sid=1>

TKDL to sign agreements with more countries, add one lakh more formulations

The Traditional Knowledge Digital Library (TKDL), the innovative tool to prevent misappropriation of traditional knowledge belonging to India at international patent offices, is in the process of expanding its access agreements with more countries and will also add one lakh more formulations of Indian streams of medicines to the existing library.

Source: <http://pharmabiz.com/NewsDetails.aspx?aid=76536&sid=1>

WHO getting ready to unveil norms on prequalification of finished pharmaceuticals

The World Health Organisation (WHO) is reviewing the draft guidelines for the scientific evaluation of finished pharmaceutical products (FPPs) by regulatory authorities which apply similarly stringent standards for quality, safety and efficacy as those recommended by it

Source: <http://pharmabiz.com/PrintArticle.aspx?aid=77137&sid=1>

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