



## **Indian Society for Clinical Research**

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### **Clinical Research in India: Patients First and Research for India -**

Theme of 9<sup>th</sup> Annual Conference of Indian Society for Clinical Research (ISCR)

**Mumbai, 23 December, 2015: Clinical Research in India: Patients First and Research for India** will be the theme of the 9<sup>th</sup> Annual Conference being held by the Indian Society for Clinical Research (ISCR) in Mumbai on January 8-9, 2016 at Hotel Trident, Bandra Kurla Complex, Mumbai. The conference, held against a much changed regulatory environment, will focus on the regulatory, ethical and operational framework for conducting clinical trials in India. With an increasing disease burden and unmet medical needs, clinical research bears a significant relevance, being an integral part of the drug development process and hence the theme of the conference is “Patients first and Research for India”.

The event will feature panel discussions, speaker series and workshops focusing on the relevance and contemporariness from the perspective of clinical operations, investigator-initiated research, accreditation, ethics, training, regulatory, pharmacovigilance, medical writing, data management, statistics and career development in clinical research. The conference will have reputed national and international speakers from across the clinical research stakeholder spectrum.

Leading up to the annual conference, four pre-conference workshops on January 7, 2016 have been scheduled for clinical researchers and students. The workshops will focus on accreditation, medical writing, risk-based monitoring and causality assessment.

For more information, please visit <http://iscr.org/home/highlights>.



## **About ISCR**

The **Indian Society for Clinical Research (ISCR)** is an association of clinical research professionals that aims to build awareness of clinical research as a specialty in India and to facilitate its growth in the country while helping to evolve the highest standards of quality and ethics. To that extent, we are fully supportive of the initiatives undertaken by regulatory authorities to create a more robust and regulated environment in India for the conduct of clinical research and will continue to work very closely with different stakeholders in the development of regulations that will safeguard and protect patients in a clinical trial.