





Clinical Trials Registry - India

National Institute of Medical Statistics ICMR, New Delhi



WHO

CTRI Bulletin Special Issue

www.ctri.in

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Launch of CTRI

20th July 2007





The Dawn of a New Era

The Clinical Trials Registry - India (CTRI), launched by Prof N.K. Ganguly, DG, ICMR on 20th July 2007, ushered in a new era in clinical research. The CTRI, hosted at the National Institute of Medical Statistics (NIMS), ICMR, New Delhi, is a free online registry of clinical trials being conducted in the country.

The registration of clinical trials in CTRI is voluntary and those conducting trials are expected to register with certain data fields before the enrolment of the first patient/participants. For details visit www.ctri.in

In this Special Issue of CTRI Bulletin, we take great pleasure in bringing to you a bird's eye view of the historic occasion.

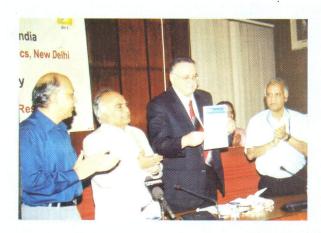


Prof. N.K. Ganguly, DG, ICMR, addressing distinguished guests

Dr. S.J. Habayeb, WHO Representative to India; Dr. Sunil Nandraj, NPO, WHO;

Dr. M. Venkateswarlu, DCG(I); Dr. S. N. Khan, DST Representative; Prof. Arvind Pandey, Director, NIMS (Administrator, CTRI); Dr. V. Muthuswamy, Sr. DDG, ICMR (Co-PI, CTRI); Prof. S.D. Seth, Chair in Clinical Pharmacology, ICMR (Advisor, CTRI);

Dr. A.N. Kapoor, DDG (Sr.), ICMR; Dr. Abha Rani Aggarwal, AD, NIMS (Coordinator, CTRI) and other scientists.



Dr. S.J. Habayeb, WHO Representative to India releasing the first CTRI Bulletin during the launch ceremony.



Prof. N.K. Ganguly joins **Dr. S.J. Habayeb**, WHO Representative to India in releasing the first CTRI Bulletin



Prof. N.K. Ganguly, "With the launch of this Registry, India will be among the few select countries like Australia, UK and the US that are making researchers accountable through registration and public disclosure of clinical trials".



Distinguished guests gathered at the Indian Council of Medical Research, New Delhi for the inauguration of CTRI.



Prof. Arvind Pandey, "We present to you a platform for registering all prospective clinical trials being conducted in India on medicinal products and information of these registered clinical trials will be made freely and publicly available on the website www.ctri.in".



Dr. S. N. Khan, DST Representative "CTRI will act as a support system for clinicians and researchers to practice transparency during the conduct of clinical trials".



Dr. S. J. Habayeb, WHO Representative to India "Congratulations to DG ICMR for the launch and establishing the Clinical Trials Registry-India. WHO will provide any assistance to CTRI and assure the future links with International Clinical Trials Registry Platform (ICTRP)".



Dr. M. Venkateswarlu, DCG(I) "The Registry is a welcome step in the regulatory process of clinical trials and any further support required from the CDSCO shall be provided to the CTRI".



Dr. Vasantha Muthuswamy, Sr. DDG, ICMR, "It is a historical occasion when the long desired initiative to launch the Registry is accomplished. All clinical trials with drugs, biotech products, vaccines, devices, biologicals and traditional medicines are expected to be registered with CTRI".

Also seen in the picture - Prof. S.D. Seth, Chair in Clinical Pharmacology, ICMR (Left) and Prof. Arvind Pandey, Director, NIMS, (ICMR) (Right).





Representatives from Academic Institutes, Pharma Industries, NGO's (PATH & DNDi), at the launch of the CTRI.



Prof. N. K. Ganguly addressing the media.

Also seen in the picture from Left to Right - Prof. S.D. Seth, Dr. V. Muthuswamy, Prof. Arvind Pandey, Prof. N.K. Ganguly, Dr. S.J. Habayeb, Dr. M Venkateswarlu.



prospectively registered before the enrollment of the first participant and to disclose details of the 20 mandatory items of the WHO International Clinical Trials Registry Platform (ICTRP) dataset. [Read more...]

The vision of the CTRI is to ensure that every clinical trial conducted in the region is prospectively registered with full disclosure of the 20-tem WHO ICTRP dataset, as well all items of the CTRI dataset, in order to 1) improve transparency and accountability, 2) improve the internal validity (details of the

trisls being conducted in India. Any researcher who plans to conduct a trial involving human participants, of any researcher who plans to conduct a trial involving human participants, of any preventive measures, lifestyte medifications, devices, educational or behavioral treatment, rehabilitation behavioral treatment, rehabilitation behavioral treatment, rehabilitation behavioral treatment of the participant. Registration is voluntary but some fields are expected of the participant. Registration in voluntary but some fields marked "WHO also need to be filled if the trial is to receive an ergostration number and fulfill WHO/CEMB consistent of the processing of the proces

Home page of CTRI (www.ctri.in)

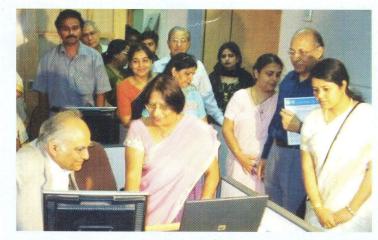
Trial Registration Set Download: (Word)[Pdf]



Left to Right - Dr. Rajat Goyal, PATH; Dr. R.H Jani, Zydus Cadila; Dr. S.K. Mishra, Dabur Research Foundation; Dr. Abha Rani Aggarwal, NIMS; Dr. Ambujam Nair Kapoor, ICMR; Prof. Prathap Tharyan, CMC, Vellore.



Prof. N.K. Ganguly, DG, ICMR, inaugurating the CTRI Unit.



Dr. Abha Rani Aggarwal, CTRI coordinator, demonstrating the process flow of www.ctri.in to DG, ICMR.



We must become the change we want to see.

-Mahatma Gandhi



National Institute of Medical Statistics, ICMR, New Delhi where CTRI is hosted.



Members of CTRI Team:

Right - Prof. Arvind Pandey, Prof. Prathap Tharyan, Prof. S.D. Seth, Dr. Abha Rani Aggarwal, Ms. Shanthi Joseph

Left - Dr. Rahmat Bano, Dr. Mohua Maulik, Ms. Anuradha Dubey, Ms. Noori Dua Back Row - Mr. Bir Singh, Mr. Harish Kumar, Mr. Anoop Upadhyay

Contact Us

Clinical Trials Registry - India

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For more information visit us at www.ctri.in

Up Coming Events

- A Regional Meeting is scheduled to be held on 12th September, 2007, at NIRRH, Mumbai.
- Journal Editors Meeting is scheduled to be held on 9th October, 2007, at ICMR, New Delhi.