



Laxmikanta Nanda <laxmikanta.nanda@gmail.com>

(LKN) Invitation to attend the second World Integrated Medicine forum, Goa, India, 23-25 January 2019

CCRH International <ccrh.international@gmail.com>

Tue, Jan 15, 2019 at 4:11 PM

To: Laxmikanta Nanda <laxmikanta.nanda@gmail.com>

Cc: Rajkumar Manchanda <rkmachanda@gmail.com>, anil23101961 <anil23101961@gmail.com>, "Dr. Deepti Singh" <dr.deepti.2821@gmail.com>

Dear Dr Nanda

It is my honor and pleasure to invite you to the second global forum on the **'Regulation of Homeopathic Medicinal Products: Advancing global collaboration'**. The Forum is a collaboration between the Central Council for Research in Homoeopathy (CCRH), an autonomous body of Ministry of AYUSH, Government of India, and the World Integrated Medicine Forum and is scheduled to be held from 23rd-25th Feb 2019 at Goa Marriot Resort & Spa, India.

This unique event builds on the success of the first World Integrated Medicine Forum held in Delhi in February 2017, and will bring together the key stakeholders involved in implementing and shaping the regulation of homeopathic medicinal products as well as manufacturers worldwide. For further details, you may like to the attached brochure, or visit: www.WIMForum.org

This exclusive, invitation only event is targeted to enable direct interaction of drug regulators/ policymakers with the manufacturers and traders of homeopathic medicinal products and will focus on the potential benefits and pitfalls of bilateral/multilateral collaboration and advance global cooperation on a synergistic basis.

Which themes will be explored?

- The challenges and benefits of regulatory collaboration between countries with a long-standing homeopathic tradition.
- How can countries with little, or a short, homeopathic tradition tap into the global regulatory community and develop appropriate national regulatory frameworks?
- Are there possibilities for bi-national and multi-national collaborations?
- How can we achieve further international convergence on minimum legal and regulatory standards?
- How can we make sure that the regulatory framework does justice to core homeopathic principles?
- How can we further advance pharmaceutical quality standards?

As a valued expert in your field, we will be honoured to have your contribution in the session on 'New homeopathic drug approval: Perspectives on setting appropriate requirements for provings and clinical data', for which we shall be registering you on complimentary basis.

Further, please note that we are currently preparing a program booklet, and as our valued contributor at the forum, we would like you to provide us with the following:

- Your Biosketch with a word count of up to 250 words for the narrative. Please include your name, main qualifications, affiliation and contact details as well. As an example, we have attached Dr. Haselen's Biosketch, for the sake of uniformity in length and contents of the submission.
- A passport size photo in a medium to high resolution JPG or other graphics format

Considering the paucity of time, please forward the biosketch and photo by reverting to this email in the NEXT FEW HOURS, please.

We will, of course, be happy to answer any further questions you may have with regard to your submission. As regards your talk-related queries, Dr. Haselen will be your go-to person.

Regards

Dr.R.K.Manchanda, M.D.(Hom), M.B.A.(Health care)

Director General,

Central Council for Research in Homoeopathy,

under Ministry of AYUSH(Ayurveda, Yoga & Naturopathy, Unani, Siddha/Sowa-Rigpa and Homoeopathy) &

2/1/2019

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Secretary, Homoeopathic Research

Liga Medicorum Homoeopathica Internationalis (LMHI)

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2 attachments



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Ministry of AYUSH,
Government of India

World Integrated Medicine Forum

Regulations of Homeopathic Medicinal Products Advancing Global Collaborations

23-25 January, 2019 | Goa, India

Organized by

Central Council for Research in Homoeopathy
Ministry of AYUSH, Govt. of India

