Date: 06.7.2018

Press Note

NIPER SMPIC seminar on : 'Impurity profiling of Pharmaceutical products: A Regulatory Requirement'

NIPER-S.A.S. has set up a dedicated Small and Medium Pharmaceutical Industry Centre (SMPIC) under the ministerial guidelines to support the Indian Pharmaceutical Induatry. The centre aims at creating commercial synergy between industry and academia. SMPIC with Indian Pharmaceutical Association (IPA) is organizing a one day seminar on 'Impurity profiling of Pharmaceutical products: A Regulatory Requirement', on 7th July, 2018. Pharmaceutical impurities are the unwanted chemicals which are introduced during the manufacturing process for drug substance, drug product, excipient or during their storage. Theses impurities should be controlled throughout the manufacturing process as these can affect safety of medicines. It is therefore, important to have an accurate detection of impurities.

This seminar shall focus on regulatory requirement of testing of impurities in pharmaceutical products, characterization of reference standards and regulatory requirements on usage of Impurity standards. Eminent speakers like Dr Jens Boertz, Pharmaceutical Product Manager, LGC -Germany, Dr Naresh Sharma, Assistant Drugs Controller CDSCO, DGHS, MoHFW (India), Mr Sidharth Sahai Malhotra, DI,CDSCO, New Delhi shall be delivering lectures on these topics.

The target audience includes personnel from the industry and researchers within the academic community. We expect about 100 participants from pharmaceutical companies, apart from academia and persons from regulatory agencies.

More information about the seminar is available at the official web-site

www.niper.gov.in or by writing to smpic@niper.ac.in

Dr Jens Boertz, Pharmaceutical Product Manager, LGC

Dr. Naresh Sharma, Assistant Drugs Controller(India),

CDSCO, DGHS, MoHFW, India

Mr Sidharth Sahai Malhotra, DI,CDSCO, New Delhi