



International Webinar

on

Ensuring safety of Medical Devices: focussed on Ocular Devices

Registration
Link

FRIDAY

April 09

03:00 PM - 05:00 PM

Speakers



Dr. V.G. Somani

Drugs Controller General of India
Central Drugs Standard Control Organization
(CDSCO), New Delhi



Dr. Rohit Saxena

Professor
All India Institute of Medical Sciences
(AIIMS), New Delhi



Dr. J Carlos Pastor

Director & Prof. of Ophthalmology
Institute of Applied Ophthalmobiology
(IOBA)



Dr. Rajeev Singh Raghuvanshi

Secretary-cum-Scientific Director
Indian Pharmacopoeia Commission (IPC),
MoHFW, Govt. of India, Ghaziabad



Dr. Promila Gupta

Principal Consultant
National Programme for Control of Blindness
& Visual Impairment (NPCBVI), New Delhi



Carmen Ruiz-Villar Fernandez-Bravo

Deputy Director, Medical Devices
Spanish Agency of Medicines &
Medical Products (AEMPS), Spain

Registration Deadline

April 08, 2021

*Seats are available

Registration Fees : 2000/- INR including GST



Indian Pharmacopoeia Commission

Ministry of Health & Family Welfare, Government of India

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OBJECTIVE

The objective of the webinar is to create awareness on safe use of ocular devices and regulatory concerns among the ocular devices manufacturers, regulators, ophthalmologists and other healthcare professionals in order to promote the safety of ocular devices manufactured and used in India.

BACKGROUND

- 01 As per Medical Device Rules, 2017 ocular medical devices are notified as drugs in India.
- 02 National Coordination Centre (NCC)- Materiovigilance Programme of India (MvPI), Indian Pharmacopoeia Commission (IPC) is authorized to monitor, record, and analyze the root cause of adverse events or risks associated with the use of medical devices and suggesting regulatory bodies for appropriate action with the sole intention of improving patient safety.
- 03 MvPI aims to promote and facilitate adverse event reporting of medical devices manufactured / used in India.

Who can attend ?

Ophthalmologists, Ocular Device Manufacturers, Distributors, Importers, Regulators, Healthcare Professionals and Researchers can attend this webinar.

OUTCOME

The outcome of this webinar is to stimulate the culture of adverse events reporting associated with the ocular devices.

Jointly organized by



Indian Pharmacopoeia Commission
(IPC)



Institute of Applied Ophthalmobiology
(IOBA)

In Collaboration with



The Spanish Agency of Medicines
and Medical Devices
(AEMPS)

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