TOXGURUKUL FOUNDATION PRESENTS

Webinar on Applied Toxicology

Title:

Formulation, route, regimen and duration of test substance administration in toxicology studies

Date: Saturday 19th Sept. 2020

Time: **8.00 pm. Indian Standard Time**

Duration: 1.5 hours (1 hr + 30 min. Q&A)

Registration link: Click here

Registration Fee: 300/- INR (for Indian nationals)/ $10 USD (for foreign nationals)

Payment link: Click here **for INR;**
Click here **for $**

Registration closing date:
Sep 17th, 2020, 6:00 PM IST with nominal fee
Sep 18th, 2020, at 12:00 pm IST with late fee (500/- INR and/or 12 USD)

Registration Procedure:

1. Register by clicking on registration link
2. You will receive a payment pending mail. Complete payment from link provided in mail or directly from payment links provided above.
3. You will receive a registration confirmation mail within 48 hours of payment

Course Moderator:
Dr.Varun Ahuja MVSc, PhD, DABT, ERT

For queries: toxgurukul.india@gmail.com

Target audience: Professionals from CROs, Pharmaceutical/Chemical/Medical device/Cosmetics/Personal care products Industry, Regulatory/Forensic/Environmental scientists, Pathologists, Academia (Faculty and research scholars e.g. MPharma, MVSc, MSc, PhD, RAs, JRFs/SRFs from any stream of pharmacological/ toxicological research).

Speaker:
Dr. Shayne Gad,
Gad Consulting Services, USA

Dr. Shayne Gad (PhD, DABT), is principal of Gad Consulting Services, a twenty-eight year old consulting firm with more than 600 clients in the US and overseas. Prior to this, he served in director-level and above positions at Searle, Synergen and Becton Dickinson, as a manager of the toxicology lab at Allied Signal, and at Chemical Hygiene Fellowship at Carnegie Mellon Institute.

Dr.Gad has more than 46 years of broad-based experience in toxicology, drug and device development, statistics, and risk assessment. A past president of the American College of Toxicology and the Roundtable of Toxicology Consultants and the recipient of the 2008 ACT Lifetime Achievement Award, Dr. Gad has authored or edited 52 books, 68 independent chapters and more than 350 papers and abstracts in the above fields. He has contributed to and has meaningful personal experience with IND (124 successfully filed to date), NDA, BLA, ANDA, 510(k), IDE, CTD, dietary supplement and PMA preparation. Dr. Gad is fully experienced with the design, conduct, analysis, and reporting of preclinical and clinical safety and pharmacokinetic studies for drugs, devices combination products and dietary supplements, and with regulatory submissions associated with them.

Few books authored by Dr. Gad include: Drug Safety Evaluation; Regulatory Toxicology; Safety Evaluation of Pharmaceuticals & Medical Devices; Statistics & Experimental design for Toxicologists; Biomaterials, Medical Devices & Combination Products; Regulations & Quality etc.

Sponsors willing to contribute financially for Student Participants, may please contact:
Dr. Mukul Pore MVSc, DABT, ERT **(President): 9822024836**
Dr. K.S. Rao MVSc, PhD, DABT **(Patron): 7337830074**