



The Conference

24 - 27 SEPTEMBER 2018

Riyadh International Convention and Exhibition Center

Safety & Risk Assessment



2030 VISION IS THE FOUNDATION ...

The Saudi Food and Drug Authority seeks to achieve the kingdom of Saudi Arabia 2030 vision by enhancing efficiency and transparency, and set up the necessary environment for the Saudi Community and business sector to endure their responsibilities. In addition, take the lead in facing challenges and grab opportunities, which will ensure the food safety, safety, quality and effectiveness of drug, and safety and efficiency of medical devices and products.

CONFERENCE OBJECTIVES

- Serves as an annual gathering connects SFDA with other stakeholders from private and public sectors.
- Exchange professional and scientific experiences with SFDA partners from the public and private sector.
- Share up-to-date best practices in regulatory sciences and other relevant disciplines.
- Transfer experiences and enhance knowledge concerning drug, food, medical devices awareness and health promotion.
- Conduct applied workshops addressing professional and scientific aspects.
- Arranging for an annual exhibition in collaboration with SFDA partners to demonstrate the updated regulatory sciences and related tools.

CONFERENCE 2018 TOPICS

SAFETY AND RISK ASSESSMENT

Food Topics

- Control and legislation of food products.
- Food safety and risk assessment.
- Animal feed and its impact on human and animal health.
- Contaminants and pesticide residues in food.
- Nutrition policies and their impact on public health.
- Legislation and specifications of innovative foods.
- Halal Food regulations.
- Halal food innovations.

Health Promotion Topics

- Evidence-based health promotion programs.
- Health education and effective communication.
- Health behavior change.

Drugs Topics

- Localization of the pharmaceutical industry.
- Medical fraud and the quality of medicines and preparations.
- Pharmacovigilance.
- Pharmacoepidemiology.
- The economics and pricing of medications.
- Control and legislation of medicines and preparations.
- Cosmetics.

Medical Devices Topics

- Clinical evaluations & investigations for medical devices.
- Medical device regulation.
- Medical devices post market surveillance and vigilance.
- Medical devices testing and calibration.
- Impact of medical device regulation.
- Cyber Security for medical devices.

ABSTRACT SUBMISSION PROCESS

Deadlines

- Abstracts submission deadline (Oral Presentations & Workshops): **25th March 2018.**
- Abstracts submission deadline (Posters): **25th July 2018.**
- All submitted abstracts will be forwarded to the Committee for review. Notifications regarding status will be sent once the review process is completed.

Abstract Submission Guideline

- Fill the abstract template which is available at www.sfdaconf.com (The format will help you to submit your abstract, therefore you are requested to follow the format carefully for your abstract submission) and fill and complete the cover page form in both Arabic and English.
- Please submit your abstract at the abstract submission center (<https://sfdaconf.com/AS/openconf.php>)
- If your abstract is accepted, you will be invited to register for the conference and pay the registration fees to confirm the presentation.



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