

*Webinar on*

# **Medical Device Regulations In The MENA Region**

*Date : 24 April 2019*

# Areas Covered

*Overview of the key areas of registration requirements for product and company approvals for medical devices in the Middle East and North Africa.*

This seminar will provide an essential overview of the key areas of requirements for product approvals for medical devices in the Middle East and North Africa.

**PRESENTED BY:**

*Ilona Putz founded PULONA Emerging Markets in 2008, based in the UAE. Her company is dedicated to creating and developing tailor-made business concepts and regulatory services for clients in the healthcare sector across the Middle East and North Africa. She holds an MBA Degree from George Washington University and attended a Finance programme at Harvard Business School.*

Date : 24 April 2019

Time : 11 : 00 AM EST

Duration : 90 Minutes

Price: \$225

# Webinar Description

This seminar will provide an essential overview of the key areas of requirements for product approvals for medical devices in the Middle East and North Africa. The programme will cover the regulatory requirements and developments in the individual countries such as Algeria, Bahrain, Egypt, Iran, Iraq, Jordan, Kuwait, Lebanon, Libya, Morocco, Oman, Palestine, Saudi Arabia, Sudan, Syria, Tunisia, UAE, and Yemen.

Gain an introduction to the medical device markets: countries, numbers, economic facts, and trends, the regulatory environment

Understand medical device regulations in the MENA region

Get an overview of registration requirements, timelines, fees

Clarify procedures for company and product registration

Medical Device regulations is an evolving era in the MENA region. Most of the MENA governments have published regulatory guidance as registration for some devices is required for importation.

# Who Should Attend ?

*Anyone involved in regulatory affairs for medical devices in the MENA region*

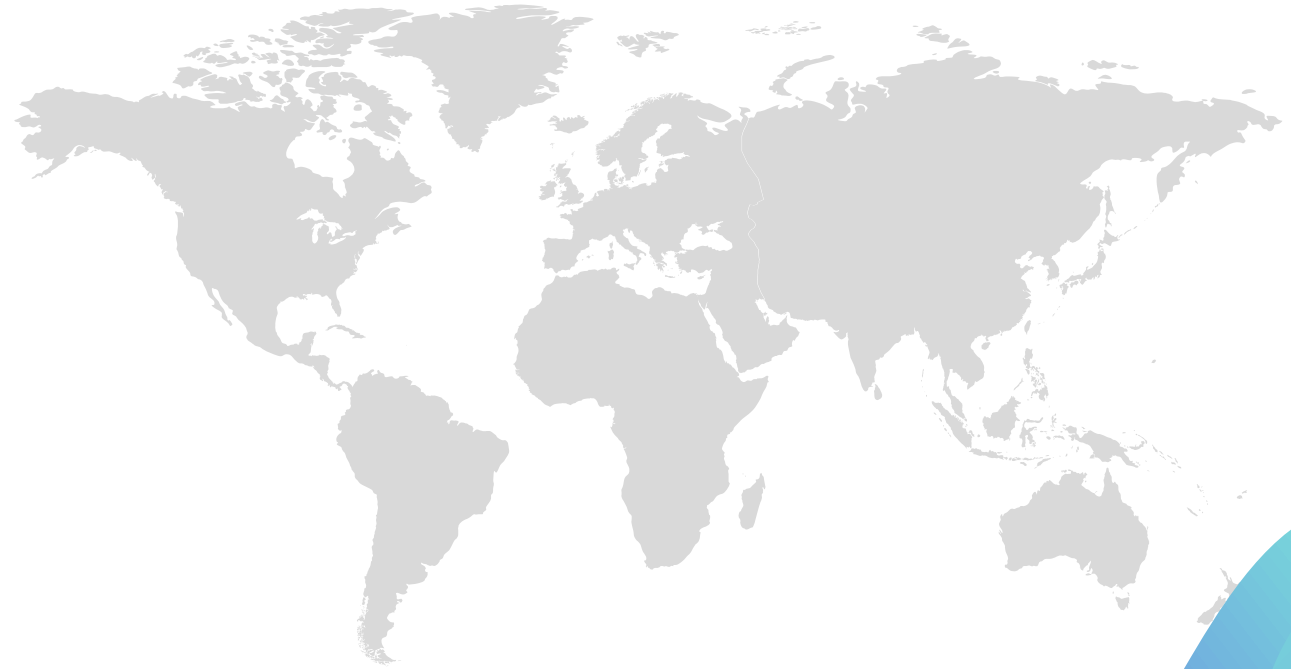
*Anyone new to the region*

*Anyone interested in an update of recent developments*



# Why Should Attend ?

*To understand the registration requirements for medical devices in the key markets in the MENA region to be able to efficiently plan submission strategies for the region.*



To register please visit:

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