GRC Online Training Hub
EDUCATORS
Axons Technology and Solutions

Webinar on

Medical Device Regulations In The MENA Region

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.

GRC Online Training Hub
EDUCATORS
Axons Technology and Solutions

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:

José Mora is a Principal Consultant specializing in Manufacturing Engineering and Quality Systems. For over 30 years he has worked in the medical device and life sciences industry specializing in manufacturing, process development, tooling, and quality systems. Prior to working full time as a consulting partner for Atzari Consulting, José served as Director of Manufacturing Engineering at Boston Scientific and as Quality Systems Manager at Stryker Orthopedics, where he introduced process performance, problem-solving, and quality system methodologies

GRC Online Training Hub EDUCATORS
Axons Technology and Solutions

On-Demand Webinar

Duration : 90 Minutes

Price: $200

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

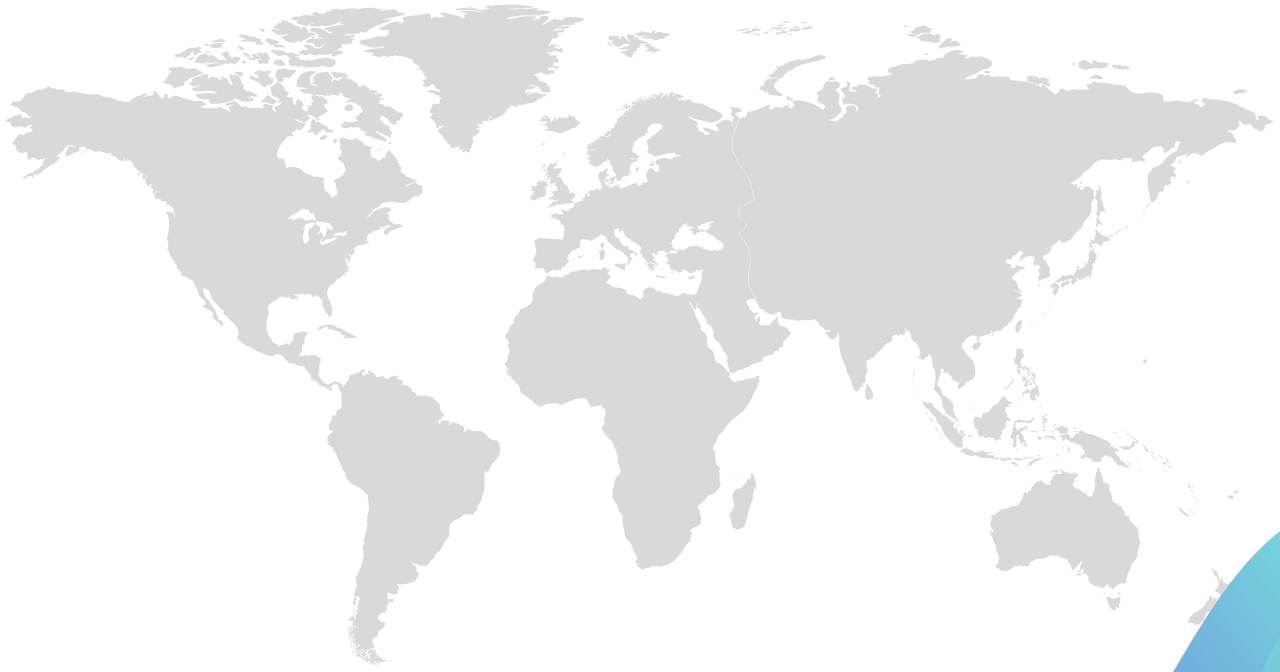
Medical Device regulations is an evolving era in the MENA region. Most of the MENA governments have published regulatory guidance as registration for some devices is required for importation.

Who Should Attend ?

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

Anyone new to the region

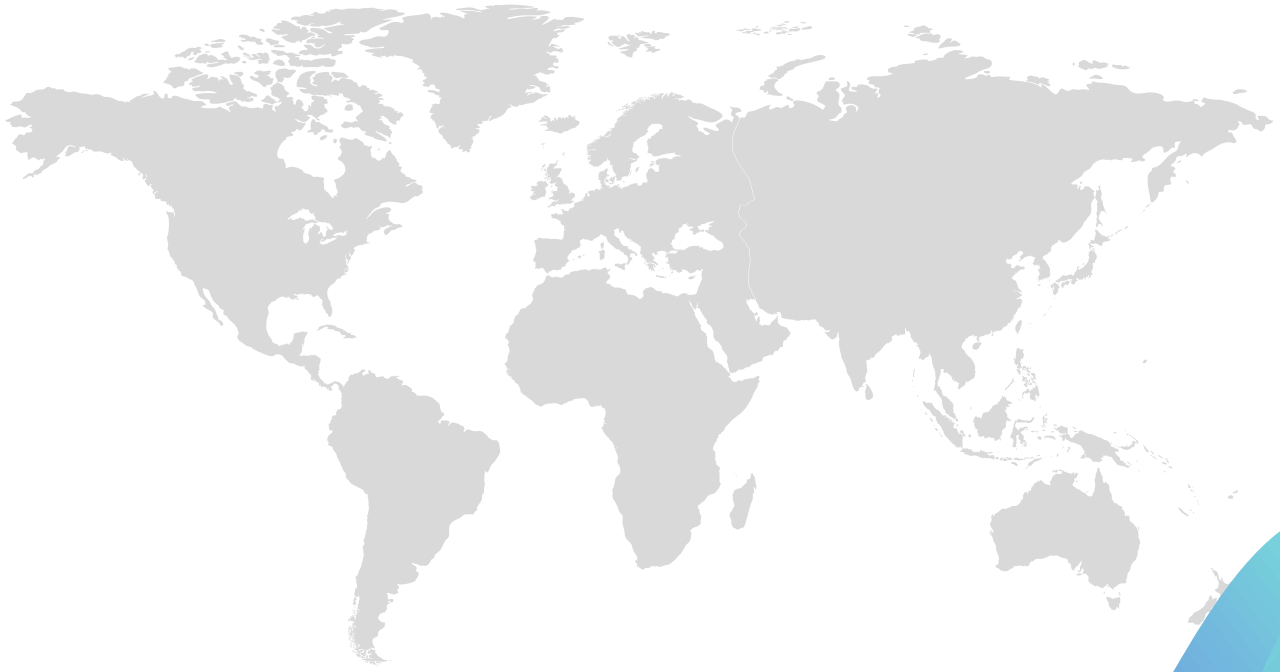
Anyone interested in an update of recent developments



GRC Online Training Hub
EDUCATORS
Axons Technology and Solutions

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.



GRC Online Training Hub
EDUCATORS
Axons Technology and Solutions

GRC Online Training Hub
EDUCATORS
Axons Technology and Solutions

To register please visit:

www.grceducators.com
support@grceducators.com
740 870 0321