

# **Middle East Pharmaceutical Contract Development & Manufacturing Organization Market Size, Share & Trends Analysis Report By Type, By Product, By Service, By Workflow, By Therapeutic Area, By End Use, By Region, And Segment Forecasts, 2025 - 2033**

<https://marketpublishers.com/r/M23D4CF07416EN.html>

Date: September 2025

Pages: 120

Price: US\$ 5,950.00 (Single User License)

ID: M23D4CF07416EN

## **Abstracts**

### **Middle East Pharmaceutical Contract Development & Manufacturing Organization Market Summary**

The Middle East pharmaceutical contract development and manufacturing organization market size was estimated at USD 3.50 billion in 2024 and is projected to reach USD 5.39 billion by 2033, growing at a CAGR of 4.90% from 2025 to 2033. The market growth is driven by increasing healthcare spending, rising prevalence of chronic diseases, and a growing demand for cost-effective drug development and manufacturing facilities.

In addition, growing government investments in healthcare infrastructure and the presence of strategic locations in the Middle East as a trade hub that connects Asia, Europe, and Africa have further contributed to market growth. Moreover, most of the established pharmaceutical companies across the globe are expanding operations in the Gulf and broader Middle East with the shifting trends of outsourcing towards contract development & manufacturing organization (CDMO) to optimize costs, accelerate time-to-market, and enhance supply chain resilience.

Moreover, rising technological advancements are enhancing both efficiency & quality within pharmaceutical development and manufacturing is expected to drive the market. Besides, the integration of automation, digital platforms, and advanced analytical tools has streamlined operations further, minimizing errors. For instance, processes such as

continuous manufacturing and single-use technologies are increasingly used to shorten production timelines and provide flexibility for both small and large molecule products. In addition, data-driven monitoring systems & real-time analytics have been implemented to ensure greater precision, compliance, and productivity across the entire value chain. Furthermore, in the Middle East, pharmaceutical innovation is driven by an expanding pipeline of biologics, biosimilars, and specialty medicines. CDMOs are stepping up to offer their expertise in complex formulations, sterile manufacturing, and the production of HPAPIs.

In addition, growing collaborative partnerships among global and emerging market players are expected to enhance the capabilities in the Middle East for biologics, cell and gene therapies, and specialized therapeutic areas. In addition, the growing expansion of R&D activities in the region and increasing requirement for outsourcing facilities due to evolving demands of the pharmaceutical pipeline are expected to drive the market over the estimated time period.

Also, the regulatory landscape in the region is more aligned with the international standards to attract investment and prioritize patient safety. Authorities such as the Saudi Food and Drug Authority & the UAE Ministry of Health and Prevention are further enhancing quality assurance and compliance frameworks. Besides, increasing requirements to harmonize regulatory guidelines with those from the U.S. FDA and EMA are underway to facilitate global market access for regionally manufactured drugs. Furthermore, growing approval processes and transparent regulatory pathways that are established to support CDMOs in maintaining competitiveness and building credibility within global pharmaceutical supply chains are expected to drive the market over the estimated time period.

## Middle East Pharmaceutical Contract Development And Manufacturing Organization Market Report Segmentation

This report forecasts revenue growth at the regional & country levels and provides an analysis of the latest industry trends in each of the sub-segments from 2021 to 2033. For this study, Grand View Research has segmented the Middle East pharmaceutical contract development and manufacturing organization market report based on type, product, service, workflow, therapeutic area, end use, and region:

Type Outlook (Revenue, USD Million, 2021 - 2033)

Small Molecule

Branded

Generic

Large Molecule

Biologics

Biosimilar

Product Outlook (Revenue, USD Million, 2021 - 2033)

API

Traditional Active Pharmaceutical Ingredient (Traditional API)

Highly Potent Active Pharmaceutical Ingredient (HP-API)

Biologics

Others

Drug Product

Oral solid dose

Semi-solid dose

Liquid dose

Others

Service Outlook (Revenue, USD Million, 2021 - 2033)

Contract Development

Pre-formulation & Formulation Development Service

Process Development & Optimization

Analytical Testing & Method Validation

Scale-up & Tech Transfer

Contract Manufacturing

API Manufacturing

Finished Drug Products Manufacturing

Packaging and Labelling

Regulatory Affairs

Others

Workflow Outlook (Revenue, USD Million, 2021 - 2033)

Clinical

Commercial

Therapeutic Area Outlook (Revenue, USD Million, 2021 - 2033)

Oncology

Infectious Diseases

Neurological Disorders

Cardiovascular Disease

Metabolic Disorders

Autoimmune Diseases

Respiratory Diseases

Ophthalmology

Gastrointestinal Disorders

Orthopedic Diseases

Dental Diseases

Others

End Use Outlook (Revenue, USD Million, 2021 - 2033)

Small Pharmaceutical Companies

Medium Pharmaceutical Companies

Large Pharmaceutical Companies

Regional Outlook (Revenue, USD Million, 2021 - 2033)

Middle East

UAE

Saudi Arabia

Kuwait

Qatar

Oman

**This report can be delivered to the clients within 3 Business Days**

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