



J C P R
THE JORDAN CENTER FOR
PHARMACEUTICAL RESEARCH
المركز الأردني للأبحاث الصيدلانية

JCPR

The Jordan Center for Pharmaceutical Research

- ✓ EMA inspected, 1st independent established CRO in MENA region (Est. 1999)

About Us

Company Overview

The Jordan Center for Pharmaceutical Research is a premier premier independent CRO delivering full-service clinical research across all phases.

We blend regional expertise with international compliance to accelerate drug development with confidence.

Founded 1999

Prof. Tawfiq Arafat



900+ Studies

Successfully completed clinical studies across varying phases.



5,000 + Volunteers

Extensive database of mixed gender gender healthy volunteers.



EMA Inspected

Among the first accredited CROs in the MENA region.



Global Reach

Regional expertise with rigorous international compliance.

OUR JOURNEY

Main Regulatory Leadership Timeline



OUR SERVICES

Bioequivalence Studies

DATA



**Study Design &
Protocol Development**



**IRB & JFDA
Submissions**



**Clinical Conduct &
Study Execution**



**Bioanalytical
Services**



**PK & Statistical
Analysis**



**Clinical Study
Report**



**Data
Management**

Clinical Facilities

FACILITIES

130 Beds

Total Bed Capacity Across 2 Sites

Dual-Site Infrastructure

JCPR operates two independent, fully equipped clinical research units. This allows high-volume simultaneous trials while maintaining absolute separation and protocol integrity.

Volunteer Safety & Comfort

Both locations are meticulously engineered to meet international clinical standards, prioritizing maximum volunteer safety, medical supervision, and comfort.



Specialized Female-Only Site Option

If required for a female-only study, one of our two sites can be fully dedicated, providing a private, segregated environment.



All-Female Staff

Staffed entirely by female medical professionals—from physicians to nurses—to respect cultural requirements.



Enhanced Sensitivity & Privacy

Strictly controlled access and separate facilities designed specifically to meet the highest standards of privacy and comfort.

Analytical Operations

LAB

CORE SYSTEMS

6 Units

LC-MS/MS Systems

High-sensitivity mass spectrometry platforms for precise bioanalysis.

EQUIPMENT MODELS

• API By SCIEX

• Shimadzu



VALIDATED ASSAYS

300+

Validated Methods



COMPLIANCE

ICH M10

Global Standards

Study Execution Process

Timeline & Phases

IVIVCLINE



COMPREHENSIVE SERVICES

Clinical Trials Phase I, III, IV

JCPR conducts comprehensive clinical trials across a wide range of therapeutic areas, providing scientifically rigorous execution from First-in-Human studies to post-marketing surveillance.

Key Therapeutic Areas Include:

Cardiovascular

Neurology

Diabetes

GI

Immunosuppressants

Oncology

Anti-infectives

Development Pipeline Capabilities

EARLY STAGE

Phase I

First-in-Human, PK/PD

CONFIRMATION

Phase III

Large Scale Safety & Efficacy, Multi-center Trials

POST-MARKET

Phase IV

Post-Marketing Surveillance, Observational Studies

Other Services



Preclinical Services



Pharmacovigilance Services

Our Services

Pharmacogenomics & Biomarker Services

Precision Medicine

Tailored Therapies



PGx-Guided Trial Design

Optimizing clinical trial protocols by incorporating pharmacogenomic inclusion/exclusion criteria to stratify patient populations effectively.



Biomarker Discovery & Validation

Comprehensive support for biomarker identification, assay development, and clinical validation to demonstrate proof of mechanism.



Genetic Testing

Advanced genotyping and sequencing capabilities, This service is provided in alliance with **GENATECHS GENATECHS Laboratories** for advanced advanced genomic and precision medicine medicine solutions .



Companion Diagnostics

Strategic support for the co-development of companion diagnostic (CDx) assays to identify patients most likely to benefit from specific therapies.



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Thank You



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