

LEADING PROVIDER OF **REGULATED SERVICES**

TO THE PHARMA INDUSTRY

DEVELOPMENT CONSULTING REGULATORY
AFFAIRS PHARMACOVIGILANCE QUALITY
MANAGEMENT & COMPLIANCE MEDICAL
DEVICE SERVICES

PharmaLex offers end-to-end regulated services that provide real value to our clients.

We differ from other contract service providers by focusing on specialized regulatory services (rather than clinical ops, manufacturing or sales support) to deliver and comply with all health agency obligations. Our global team has more than 850 local knowledge experts.

PharmaLex supports pharma companies throughout the entire product lifecycle, ensuring compliance with pharmaceutical regulations and providing vital scientific and strategic advice on drug development. Our services extend beyond market approval and include product maintenance post launch activities.

OUR SERVICES

DEVELOPMENT CONSULTING

Scientific, regulatory and strategic advice for clients with development-stage drugs, including scientific writing and project management, as well as medical affairs services.



REGULATORY AFFAIRS

Support for all activities and processes required to achieve and maintain compliance with pharmaceutical regulations and laws.



PHARMACOVIGILANCE

Support or full management of drug adverse event monitoring and reporting required to comply with regulations; pharmacoepidemiology studies.



QUALITY MANAGEMENT & COMPLIANCE

Guidance and support for establishing and maintaining quality control systems in compliance with regulations for pharmaceuticals.



MEDICAL DEVICE SERVICES

Regulatory affairs and quality management & compliance services for medical devices.



PROVIDING SERVICES EXACTLY WHERE OUR CLIENTS NEED US

NORTH AMERICA

Regional hubs
USA

LATIN AMERICA

Regional hubs
BRAZIL
PUERTO RICO

MIDDLE EAST

Regional hubs

AFRICA

Regional hubs
CHINA
INDIA

CIS & RUSSIA

Regional hubs

EUROPE

Regional hubs

BELGIUM
BULGARIA
FRANCE
GERMANY
IRELAND
ITALY
LITHUANIA
NORDIC REGION
SPAIN
SWITZERLAND
UK

ASIA

Regional hubs
CHINA
INDIA



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