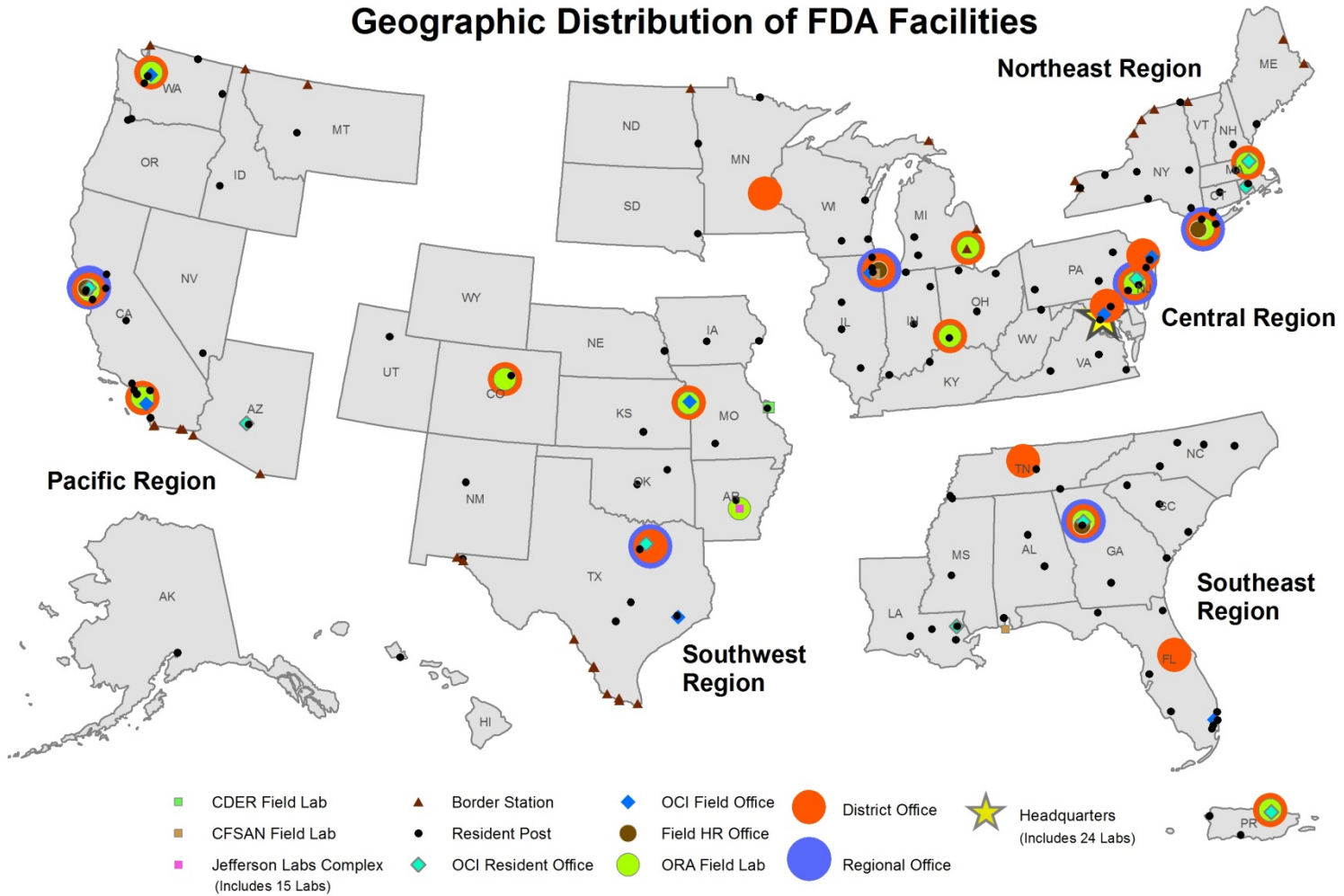


GEOGRAPHICAL DISTRIBUTION OF FDA FACILITIES



Geographic Distribution of FDA Facilities



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GLOSSARY OF ACRONYMS

3D	3-Dimensional
ACOMS	Advisory Committee Oversight and Management Staff
ACSI	American Customer Satisfaction Index
ADE	Adverse Drug Experience
ADEPT	Autonomous Diagnostics to Enable Prevention and Therapeutics
ADHD	Attention-Deficit / Hyperactivity Disorder
ADUFA	Animal Drug User Fee Act
AGDUFA	Animal Generic Drug User Fee Act
AMP	Real Property Asset Management Plan
ANDA	Abbreviated New Drug Application
ANPRM	Advance Notice of Proposed Rulemaking
APEC	American Customer Satisfaction Index
ARL	Arkansas Regional Laboratory
ARS	Agriculture Research Service
ARS	Acute Radiation Syndrome
B&F	Buildings and Facilities
BA	Budget Authority
BACPAK	Bacterial Pathogen Knowledge Base
BARDA	Biomedical Advanced Research and Development Authority
BIMO	Bioresearch Monitoring
BLA	Biologic License Application
BMAR	Backlog of Maintenance and Repairs
BPA	Bisphenol A
BPCA	Best Pharmaceuticals for Children Act
BRF	Beltsville Research Facility
BsUFA	Biosimilars User Fee Act
CBER	Center for Biologics Evaluation and Research
CBP	Customs and Border Protection
CBRN	Chemical, Biological, Radiological, and Nuclear

CDC	Centers for Disease Control and Prevention
CDER	Center for Drug Evaluation and Research
CDRH	Center for Devices and Radiological Health
CERSIs	Centers of Excellence in Regulatory Science and Innovation
CFR	Code of Federal Regulations
CFSAN	Center for Food Safety and Applied Nutrition
cGMP	current Good Manufacturing Practice
CIADM	Centers for Innovation in Advanced Development and Manufacturing
CIO	Chief Information Officer
CMS	Centers for Medicare & Medicaid Services
CMV	Cytomegalovirus
CORE	Coordinated Outbreak Response and Evaluation
CORES	Collaborative Opportunities for Research Excellence in Science
CRADA	Cooperative Research & Development Agreement (CRADA)
CSU	Central Shared Use
CT	Computed Tomography Imaging
CTP	Center for Tobacco Products
CUP	Central Utility Plant
CVM	Center for Veterinary Medicine
CY	Calendar Year
DARPA	Defense Advanced Research Projects Agency
DHRD	Division of Human Resource Development
DHS	Department of Homeland Security
DILI	Drug-Induced Liver Injury
DIO	Division of Import Operations
DNA	DeoxyriboNucleic Acid
DOD	Department of Defense
DSC	Drug Safety Communication
DTC	Direct-To-Consumer
DTRA	Defense Threat Reduction Agency
DxOD	Diagnostics on Demand

EADB	Estrogenic Activity Database
EDKB	Endocrine Disruptor Knowledge Base
EDR	Electronic Data Room
EDSR	Electronic Document Submission and Review
EIR	Entrepreneurs in Residence
EMA	Economically Motivated Adulteration
eMDR	Electronic Medical Device Reporting
E.O.	Executive Order
EON	Emergency Operations Network
EON IMS	Emergency Operations Network Incident Management System
ESPC	Energy Savings Performance Contract
ESRD	End-Stage Renal Disease
ETASU	Elements to Assure Safe Use
EUA	Emergency Use Authorizations
FACA	Federal Advisory Committee Act
FAERS	FDA Adverse Event Reporting System
FAO	Food and Agriculture Organization
FATA	Federal Anti-Tampering Act
FCC	Forensic Chemistry Center
FCI	Facility Condition Index
FCN	Food Contact Substance Notification
FD&C Act	Federal Food, Drug and Cosmetic Act
FDA	Food and Drug Administration
FDAAA	Food and Drug Administration Amendments Act of 2007
FDAMA	Food and Drug Administration Modernization Act
FDASIA	Food and Drug Administration Safety and Innovation Act
FDA-TRACK	FDA-wide performance management system
FDCA	Federal Food, Drug and Cosmetic Act
FEMP	Federal Energy Management Program
FERN	Food Emergency Response Network
FFDM	Full-Field Digital Mammography
FMT	Fecal Microbiota Transplantation

FOI	Freedom of Information Act
FOIA	Freedom of Information Act
FPC	Federal Partners Collaboration
FSIS	Food Safety Inspection Service
FSMA	Food Safety Modernization Act
FSVP	Foreign Supplier Verification Programs
FTE	Full Time Equivalent
FVM	Foods and Veterinary Medicine
FY	Fiscal Year
GDUFA	Generic Drug User Fee Amendments
GFI	Guidance for Industry
GIS	Geographic Information System
GMP	Good Manufacturing Practices
GO	Global Regulatory Operations and Policy Directorate
GSA	General Services Administration
GUDID	Global UDI Database
HDE	Humanitarian Device Exemption
HHS	Department of Health and Human Services
HIV	Human Immunodeficiency Virus
HQ	FDA Headquarters
HRWG	High Risk Working Group
HSP	Human Subject Protection
HUD	Humanitarian Use Device
HVAC	Heating, Ventilation, and Air Conditioning
ICCR	International Cooperation on Cosmetics Regulation
ICH	International Conference on Harmonization
ICOR	International Consortium of Orthopedic Registries
IDE	Investigational Device Exemption
IFT	Institute of Food Technologists
IMDRF	International Medical Device Regulators Forum

IND	Investigational New Drug
IOM	Institute of Medicine
IRB	Institutional Review Board
IT	Information Technology
ITACS	Import Trade Auxiliary Communications System
IVD	In Vitro Diagnostics
JLC	Jefferson Labs Complex
LSBC	Life Sciences-Biodefense Laboratory Complex
MAQC	MicroArray Quality Control
MCM	Medical Countermeasure
MCMi	Medical Countermeasures initiative
MDE	Medical Device Epidemiology
MDIC	Medical Device Innovation Consortium
MDR	Medical Device Reporting
MDSAP	Medical Device Single Audit Program
MDSP	Medical Device Shortages Program
MDUFA	Medical Device User Fee Amendments
MDUFMA	Medical Device User Fee and Modernization Act
MERS	Middle East Respiratory Syndrome
MERS-CoV	Middle East Respiratory Syndrome Coronavirus
MFRPS	Manufactured Food Regulatory Program Standards
microRNA	Micro Ribonucleic Acid
MIT/HST	Massachusetts Institute of Technology/Health Science and Technology
MOD	Module
MQSA	Mammography Quality Standards Act
MRI	Magnetic Resonance Imaging
M RTP	Modified Risk Tobacco Product
NA	Not Approvable
NADA	New Animal Drug Application

NARMS	National Antimicrobial Resistance Monitoring System
NCBI	National Center for Biotechnology Information
NCTR	National Center for Toxicological Research
NDA	New Drug Application
NGO	Non-governmental Organization
NIH	National Institutes of Health
NIOSH	National Institute for Occupational Safety and Health
NME	New Molecular Entity
NSABB	National Science Advisory Board for Biosecurity
NSAID	Non-Steroidal Anti-Inflammatory Drugs
NSE	Not Substantially Equivalent
NYTS	National Youth Tobacco Survey
OBE	Office of Biostatistics and Epidemiology, CBER
OC	Office of the Commissioner
OCAC	Office of Cosmetics and Colors
OCC	Office of the Chief Counsel
OCE	Office of Compliance and Enforcement
OCET	Office of Counterterrorism and Emerging Threats
OCI	Office of Criminal Investigations
OCM	Office of Crisis Management
OCP	Office of Combination Products
OCS	Office of the Chief Scientist
OCT	Optical Coherence Tomography
OCTC	Office of the Counselor to the Commissioner
OEA	Office of External Affairs
OECD	Organization for Economic Co-Operation and Development
OFVM	Office of Foods and Veterinary Medicine
OGCP	Office of Good Clinical Practice
OGROP	Office of Global Regulatory Operations and Policy
OHCA	Office of Health and Constituent Affairs
OIM	Office of Information Management
OIP	Office of International Programs

OIR	Office of In Vitro Diagnostics and Radiological Health
OL	Office of Legislation
OMA	Office of Media Affairs
OMB	Office of Management and Budget
OMPT	Office of Medical Products and Tobacco
OO	Office of Operations
OOPD	Office of Orphan Products Development
OPP	Office of Policy and Planning
OPT	Office of Pediatric Therapeutics
ORA	Office of Regulatory Affairs
ORISE	Oak Ridge Institute for Science and Education
ORRR	Other Rent and Rent Related
ORSI	Office of Regulatory Science and Innovation
OSE	Office of Surveillance and Epidemiology, CDER
OSI	Office of Scientific Integrity
OSMP	Office of Special Medical Programs
OSPD	Office of Scientific and Professional Development
OTC	Over-the-counter
PAC	Pediatric Advisory Committee
PAD	Program Activity Data
PAHO	Pan American Health Organization
PAHPRA	Pandemic and All-Hazards Preparedness Reauthorization Act of 2013
PATH	Population Assessment of Tobacco and Health
PB	President's Budget
PC	Preventive Control
PDC	Pediatric Device Consortia
PDMA	Prescription Drug Marketing Act
PDUFA	Prescription Drug User Fee Act
PHEMCE	Public Health and Emergency Countermeasures Enterprise
PIC/S	Pharmaceutical Inspection Convention and Cooperation Scheme
PMA	Premarket Approval Application
PREA	Pediatric Research Equity Act

PREDICT	Predictive Risk-Based Evaluation for Dynamic Import Compliance Targeting
PRISM	Post-Licensure Rapid Immunization Safety Monitoring
PTN	Pediatric Trials Network
QSDAR	Quantitative Spectroscopic Data-Activity Relationships
REMS	Risk Evaluation and Mitigation Strategy
RFCTG	Regulators Forum Cell Therapy Group
RTA	Refusal to Accept
SE	Substantially Equivalent (when used by Device and Biologics Programs)
SEQC	Sequencing Quality Control
SE	Substantial Equivalence
SLEP	Shelf Life Extension Program
SEQC	Sequencing Quality Control
SNS	Strategic National Stockpile
SLEP	Shelf Life Extension Program
SP	Strategic Priority
SRL	Southeast Regional Laboratory
SW	Southwest
TB	Tuberculosis
TCORS	Tobacco Centers of Regulatory Science
TB	Tuberculosis
TIMS	Tobacco Inspection Management System
TPMP	Tobacco Product Manufacturing Practice
TPSAC	Tobacco Product Scientific Advisory Committee
UDI	Unique Device Identification
UESC	Utility Energy Service Contract
UF	User Fee
UESC	Utility Energy Service Contract
UN	United Nations

USAMRIID	United States Army Medical Research Institute for Infectious Diseases
USC	United States Code
USDA	United States Department of Agriculture
USP	U.S. Pharmacopoeia
VAERS	Vaccine Adverse Event Reporting System
VICH	Veterinary International Conference on Harmonization
VKA	Vitamin K Antagonist
WD	Withdrawn
WEAC	Winchester Engineering and Analytical Center
WHO	World Health Organization

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