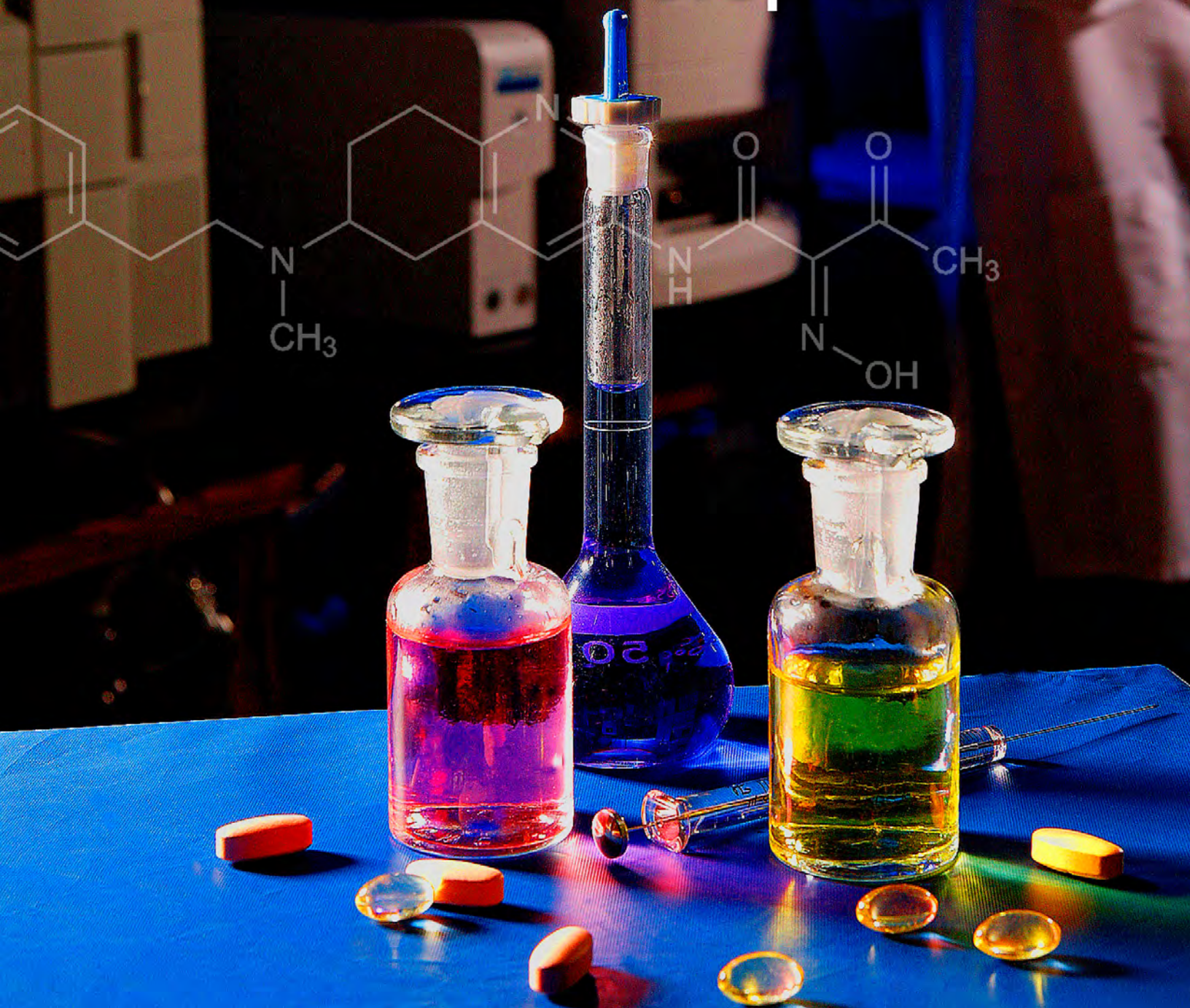


# Integrated Pharmaceutical Development



SOUTHWEST RESEARCH INSTITUTE®



**S**outhwest Research Institute® (SwRI®) is a nonprofit contract research and development organization offering comprehensive services spanning drug design and discovery through Current Good Manufacturing Practice (cGMP) drug substance and drug product supply for clinical trials. Integrated program management, consulting, technical writing, and preparation of regulatory applications services are also available.

Drug  
Discovery

Biochemistry

Drug  
Substance  
Development

Drug  
Product  
Development

cGMP  
Manufacturing

Program  
Management  
& Regulatory  
Submissions

## Southwest Research Institute®

Founded in 1947 as an independent, nonprofit research and development organization, Southwest Research Institute provides significant research, engineering, and evaluation resources for industry, business, and government. With nine technical divisions and state-of-the-art laboratories, the Institute uses a multidisciplinary, integrated approach to solving complex problems in science and applied technology. Subject to the client's wishes, programs are kept confidential. As part of a long-held tradition, patent rights arising from sponsored research at the Institute are often assigned to the client. SwRI generally retains the rights to Institute-funded advancements.

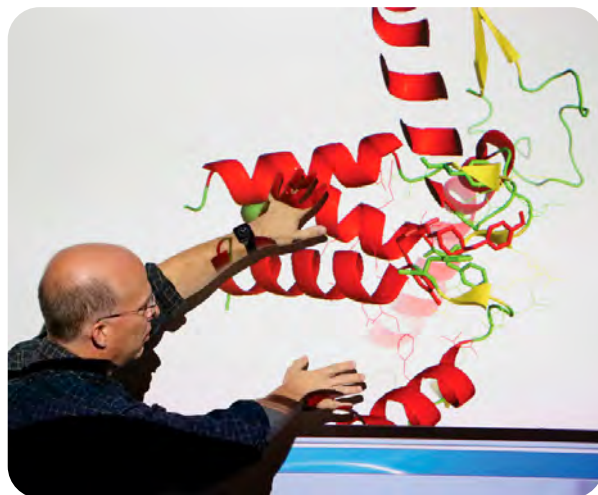
©2019 Southwest Research Institute.  
All rights reserved.

Southwest Research Institute and SwRI are registered trademarks in the U.S. Patent and Trademark Office.

### Drug Discovery

Biostructure-based drug design with proprietary Rhodium™ docking platform

- QSAR services — pharmacophore modeling, PCA, homology models
- Screening library synthesis
- *In vitro* assay development
- Hit to lead optimization
- Physicochemical properties determination and salt screening
- Scale-up for *in vivo* studies

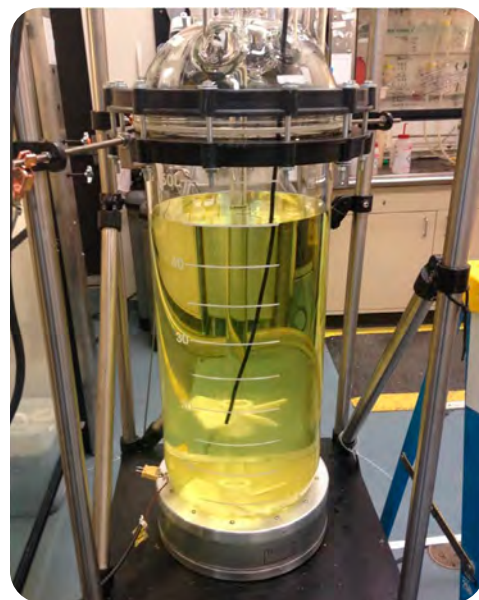


### Biochemistry

- Tissue culture, microbiology, molecular biology, antibodies
- Protein expression, purification, characterization
- Protein engineering (activity, stability, stereoselectivity)
- Enzyme kinetics, calorimetry, UV/Vis/fluorescent spectroscopy
- Cell based assays, ELISA
- Viral expression and characterization

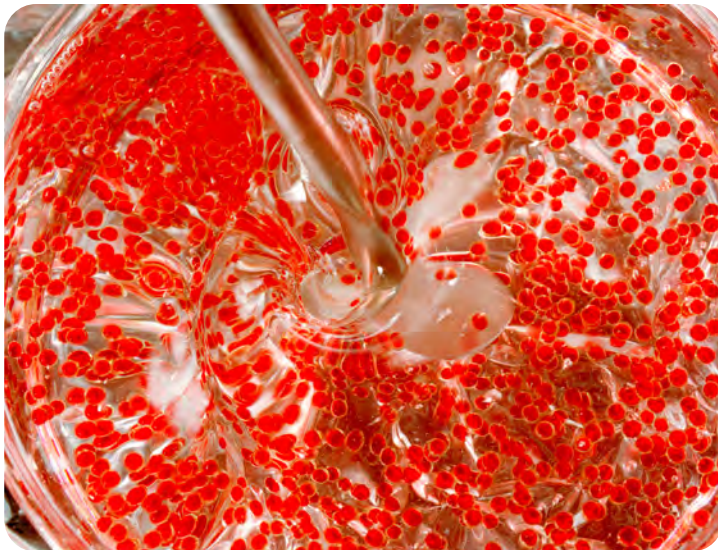
### Drug Substance (API)

- Technology transfer and evaluation of existing process
- *De novo* process development
- Isolation and characterization of process impurities
- Process optimization and parametric studies (design space)
- High potency drugs
- Controlled substances — schedule II-V
- Specialized needs (chromatography, flow chemistry, others)
- Physicochemical properties and salt screening
- Analytical methods development and qualification or validation
- ICH stability studies
- Forced degradation
- Reference standard qualification
- Specifications development



## Drug Product (Formulation)

- Pre-formulation/formulation development for small and large molecules, controlled substances, potent compounds, toxins
- Oral, parenteral dosage forms
- Scale-up and technology transfer
- Microencapsulation — extrusion, fluidized bed, spray drying, liposomes, micelles
- Milling (hammer mill, ball mill, jet mill, bead mill)
- Granulation (fluidized bed coater, spray dryer, pan coater, extrusion)
- Emulsion/suspension (homogenizer, polymerization, pickering emulsion)
- Specialized dosage forms (implantable delivery systems, sustained release, particle size reduction, protein conjugates and targeted delivery systems)
- Particle engineering — granulation, emulsions, suspensions, electrospinning, core shell particles, polymerization, morphology design
- Engineering and consulting – custom equipment design and demonstration
- Analytical methods development and validation/qualification



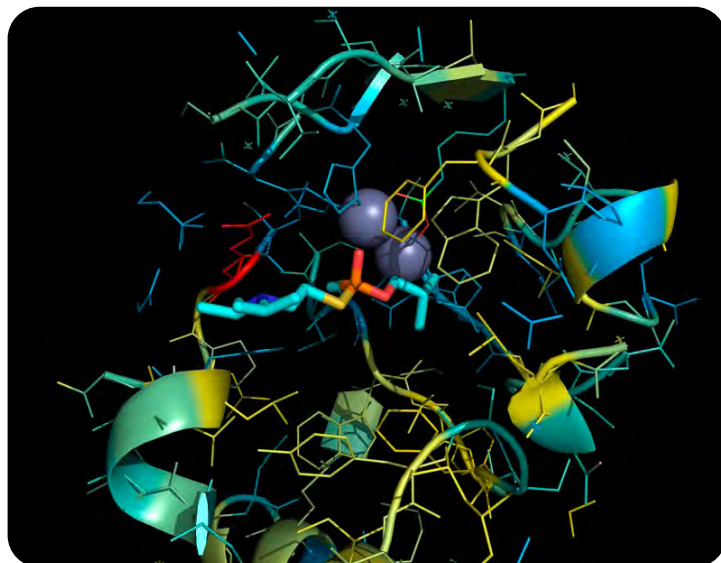
## cGMP Manufacturing (Drug Substance and Drug Product)

- Gram to kilogram scale manufacturing (drug substance, drug product)
- Technology transfer
- FDA inspected facility
- ISO 9001:2008 quality system
- cGMP chromatography services
- High potency and controlled substances
- Specialized processes
- Release testing for drug substance and drug product



## Rhodium™ — New Dimension in Protein Docking

- Unbiased comprehensive *in silico* docking on protein structures
- Extremely high throughput
- Demonstrated efficacy in multiple druggable targets
- Superior selectivity for selection of true ligands
- Superior accuracy in predicting binding poses
- Ideal for allosteric sites and protein-protein interactions
- Available as a service — no contracts or licenses required







*Southwest Research Institute® is a premier independent, nonprofit research and development organization. With eleven technical divisions, we offer multidisciplinary services leveraging advanced science and applied technologies. Since 1947, we have provided solutions for some of the world's most challenging scientific and engineering problems clients.*

**We welcome your inquiries.  
For additional information, please contact:**



Advanced science. Applied technology.

Darrel Johnston  
Director  
(210) 522-2160  
[darrel.johnston@swri.org](mailto:darrel.johnston@swri.org)

Pharmaceuticals and Bioengineering Department  
Chemistry and Chemical Engineering Division  
Southwest Research Institute  
6220 Culebra Road • P.O. Drawer 28510  
San Antonio, Texas 78228-0510



[pharmdev.swri.org](http://pharmdev.swri.org)



The Chemistry and Chemical Engineering Division of Southwest Research Institute has achieved certification to ISO 9001:2015, an internationally recognized quality standard.

**Find us on**



**[swri.org](http://swri.org)**

San Antonio, Texas • (210) 522-2122 • [ask@swri.org](mailto:ask@swri.org)

Designed & printed by SwRI MPS