

## API DEVELOPMENT & MANUFACTURING

### Exclusively Manufactured in the USA

- Qualified** - State-of-the-art equipment & facilities
- Quality** - Highest Purity Drug Ingredients
- Value** - Consistent, respected, reliable
- Validated** - FDA registered and inspected
- Trusted** - Most stringent quality system
- Support** - DMF submission
- Security** - Made in the USA



### API Focus & Expertise

- Small Volume Support
- Small Molecule Synthesis
- Orphan Drug Substance Quantities
- Chlorinations / Chloride Compounds
- Salts of Complex Organic Compounds
- Inorganic & Boutique Organic Synthesis
- Full Compliance vs. Atypical Standards
- Low Bioburden, Low Endotoxin Demands
- Parenteral / Oral / Transdermal Applications



# API Support Package

## Process Development – Manufacture of Registration Batches – Ongoing Support

### Analytical Support

- Analytical Method Validation
- Transfer of Analytical Methods
- Custom Analytical Methods and Specifications
- Bioburden and Endotoxin Testing
- Complete Impurity Profile
- Elemental Impurities
- Residual Solvents



### Development Support

- Stability Study
- Custom GMP Services as needed
- Custom Labeling and Packaging
- Manufacture of API Registration Batches
- Drug Master File submission
- Letter of Authorization
- Efficient Development Timeline
- High-touch management of your project

### Ongoing Support - Post FDA Approval

- Commercial Manufacturing of your API
- On Site Audits
- Annual Product Review
- Management of Change
- Post Submission Change Notification
- Full support through the life of your product

