



Stason Pharmaceuticals, Inc.
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Vital Statistics

Year Founded: 1994

Number of Employees: 60

Who We Are

STASON PHARMACEUTICALS, INC. is a vertically integrated cGMP contract development organization that provides complete turn-key drug development services for oral products. Our technology platforms include immediate & delayed release tablets and capsules, and fast oral disintegrating tablets. We provide our clients with flexibility, from bench top to commercial manufacturing and have facilities throughout the US, and Japan, China and Taiwan.

Services Offered

- Pre-Formulation & Formulation Development
- Method Development & Validation
- cGMP Manufacturing of preclinical through phase 3 studies
- Quality Control & Stability Studies
- Scale-up & technology transfer
- Pilot to commercial production
- High Containment Operations
- Packaging, Storage & Distribution
- Regulatory Services (IND, ANDA, 505 (b)(2), NDA)

- DEA Analytical Permit for CII-CIV
- 467 Compliance assistance

Facilities

Stason offers both Conventional Operations as well as High Containment Operations.

The manufacturing facility is 36,000 square feet, Class 100,000. Our capacity is 0.5kg – 600kg and can produce over 2 billion oral doses per year.

Stason currently manufactures APIs for world-wide clients in it's FDA inspected High Containment facilities. We currently produce high potency APIs and finished products at commercial scale. We have the capability to handle Category I-IV compounds (0.03µg/m³).

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