Selecting the right services for your new compound

From planning to filling to tracking, Vetter and Sentry can guide your compound through the clinical trial manufacturing process with quality, attention to detail, and efficiency.

Vetter Development Service

Clinical manufacturing starts with Vetter

Aseptic filling and lyophilization for early clinical trial phases

Services

- Formulation support
- Process development
- Clinical trial manufacturing of vials and syringes
- Analytical services
- Regulatory support

Capabilities

- Automated vial and syringe filling lines in restricted access barrier systems (RABS), near Chicago
- Lyophilization
- Facility and processes designed to maximize production yield with provided API
- QA testing in adherence with FDA, EMA, and a wide range of international standards

Sentry BioPharma Services

Sentry streamlines the path to clinic

Packaging services and supply chain management services

Services

- Cold chain production management activities
- cGMP storage (ambient to -90°C)
- Labeling, kitting, and secondary packaging
- Shipping services
- Product return and destruction programs
- Import/export optimization

Capabilities

- Highly protected risk (HPR) 50,000 ft^2 facility within the Indianapolis International Airport's Foreign Trade Zone, which allows products from outside the US to be held at temperature pending regulatory approval for importation from the FDA, CBP, and/or DEA
- Verified-Accredited Wholesale Distributors® (VAWD) facility
- Compliant with 21 CFR (parts 210 and 211)
- Licensure from all 50 US states and current inspections from many regulatory bodies

Both Vetter and Sentry offer their services in a high quality manner throughout the entire clinical manufacturing process along with the expertise you need to navigate the critical clinical manufacturing process.

VETTER
Answers that work
www.vetter-pharma.com

For US inquiries, please contact infoUS@vetter-pharma.com

Expertise throughout your compound´s clinical journey