Simtra
BioPharma Solutions

Made for this

OUR CAPABILITIES

Your Partner for Vital Injectable Products

With 65+ years focused exclusively on sterile injectables, Simtra BioPharma Solutions brings integrated scientific, technical and manufacturing expertise — from development through global commercial fill/finish. As an agile, independent contract development and manufacturing organization (CDMO), we partner with pharma and biotech companies to reduce risk, scale with confidence and accelerate delivery of high-quality injectables to patients.

A Commitment to Precision, Excellence & Growth

Scientific & technical expertise

- Full-service continuum — development through commercial fill/finish
- 150M+ units produced annually
- Lyophilization Center of Excellence

Operational performance

- Partner to 50%+ of top 20 pharma companies
- World-class US and EU facilities
- Committed to delivering right first time, on time and in full

Regulatory support & quality assurance

- 65+ audits per year
- Inspected by 19+ regulatory agencies globally
- Products marketed into 120+ countries
- Robust quality assurance and quality control at every step

Future-focused capabilities

- Ongoing investment in global operations
- US & EU expansion of services and capacity

From Lab Through Launch

Integrated services minimize risk, avoid handoffs and safeguard product knowledge — helping programs move faster and stay on track.

Singular Focus on Sterile Injectables

End-to-end expertise means proven processes and the ability to deliver even the most complex sterile injectables efficiently and cost-effectively.

Delivery platforms

Prefilled syringes | Cartridges | Liquid vials | Lyophilized vials

Product types

- Cytotoxics
- Bioconjugates, including antibody-drug conjugates (ADCs)
- Highly potent compounds
- Biologics
- Small molecules
- Diluent syringes

Global capabilities support standard and complex sterile injectable products.

At Simtra, we're *made for this.* Our proven sterile injectables expertise and integrated capabilities make us a full-service, agile partner — ready to scale and bring your project to market with confidence.

- Integrated, co-located preclinical development through commercial launch services
- Meets FDA, JP and EU regulatory standards
- EHS certification according to ISO 14001
- Automated and manual packaging options
- Qualified cold chain storage: -70 °C / -20 °C / 2–8 °C / 15–25 °C

~2500 employees worldwide

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