Pfizer CentreOne

What does it take to deliver with exact SI-ence?

Navigating the intricate sterile injectable (SI) landscape, supporting the unique needs of your project and delivering high quality demands an exact SI-ence.

An altogether different perspective

Pfizer CentreOne is **here to help.**

In a recent podcast with Pharmaceutical Technology, our experts Dr. Martin Gonzales, Senior Manager of Product Development at Pfizer, and Katharine Sparhawk, Senior Manager of Business Development at Pfizer CentreOne, discuss what it takes to support your SI therapy on its journey to patients effectively.

Listen to the podcast

How does Pfizer CentreOne support the production of your SI drug products?

As a CDMO within the Pfizer network, we offer comprehensive support for the production of your SI product. Harnessing deep scientific and technical expertise, combined with cutting-edge facilities, we can provide you with:

- **Cutting-edge development and manufacturing solutions:**
  We handle both small and large-scale development and commercial manufacturing of your SI.
- **Diverse dosage forms:**
  Our centers of excellence specialize in producing liquid, lyophilized, pre-filled syringes, vials, and cartridges.
- **Adaptable production:**
  With multiple filling lines and diverse packaging options, we offer flexibility across various vial, cartridge, and syringe sizes.

What communication strategies and tools do you use to align expectations during tech transfer and project progression?

- By implementing clear points of contact and designating decision-makers, we're able to streamline the tech transfer processes and prevent bottlenecks.
- At the end of each project, we identify and implement any key learnings to drive continuous improvement.
- We utilize tools such as electronic batch records, co-develop analytical methods between teams, and perform regular equipment and software updates to ensure seamless regulatory compliance.

How do you ensure flexibility and transparency in capacity planning for SI projects?

Through open communication, we align our network's capabilities with your goals.

We work with our site teams to develop both short-term and long-term forecast planning, ensuring we can fulfill your immediate needs and plan for the next five years.

Your project is prioritized, secured and integrated into our everyday processes, supported by a dedicated team with direct communication channels.

Can you ensure consistent quality and compliance with regulatory standards?

With over 80 years of experience in the regulatory field, our team is equipped to provide you with end-to-end regulatory services covering more than 150 countries.

We'll collaborate closely with you, guiding you through the regulatory landscape from development and submission to commercialization.

Ready to start your SI journey?

Discover how we can help you bring your high-quality therapies to patients with exact SI-ence as an altogether different CDMO.

Learn more