Singota Solutions is a contract development and manufacturing organization (CDMO) focused on moving products through the drug development pipeline faster by being agile, accountable, and transparent.

Development & Quality Control

DEVELOPMENT & GMP TESTING LABORATORY

Our on-site quality control, development, and testing capabilities at Singota span a wide range, allowing us to meet specific client needs in a streamlined and efficient manner. Services include analytical development, process development, and analytical testing. Early phase development support upstream of manufacturing includes material compatibility studies, forced degradation, and method transfers/validation. Post manufacturing support includes release testing, stability, testing, and COA generation.

Manufacturing

ASEPTIC FILLING & FINISHING

Our gloveless, robotic, aseptic filling workcell provides a state-of-the-art filling process for manufacturing injectable therapies. Singota specializes in filling small batch parenterals into ready-to-use vials, syringes, or cartridges. Finishing services include customized labeling, kitting, and secondary packaging with the ability to handle temperature sensitive materials and clinical trial requirements.

Supply Chain

SCM & COLD CHAIN

Singota manages a wide range of materials from APIs and excipients to finished products. We have the capability to handle toxic, potent, flammable, and hazardous materials. Our secure, cGMP warehouse offers storage conditions including controlled room temperature (15-25°), 2-8°C, -20°, and -80°C. Singota also provides distribution of clinical trial materials, as well as cold chain solutions for temperature sensitive products.

Contact us for your project:
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Focused on Faster

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