

SCALABLE TALENT FOR GMP-COMPLIANT MANUFACTURING

TALENT + TECHNOLOGY SOLUTIONS FOR PHARMACEUTICAL MSAT & PRODUCTION EXCELLENCE

As biotech and pharmaceutical organizations bring therapies from the lab bench to large-scale production, operational excellence and compliance are paramount. From aseptic fill-finish to cell and gene therapy manufacturing, companies must balance speed, quality, and regulatory rigor.

Dexian is your partner for building end-to-end pharmaceutical manufacturing teams. We deliver specialized MSAT, quality, and operations talent across sterile manufacturing, tech transfer, and commercial-scale production—ensuring your products reach patients safely and efficiently.

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Expertise Across Pharmaceutical Manufacturing Lifecycles



ASEPTIC & STERILE PROCESSING

- Aseptic Manufacturing Operators
- Sterile Filling & Cleanroom Technicians
- Microbiology & Environmental Monitoring Specialists
- Gowning & Containment Protocol Experts



CELL & GENE THERAPY PRODUCTION

- Cell Therapy Manufacturing Technicians
- Viral Vector Production Scientists
- Bioprocess Engineers (Upstream & Downstream)
- Single-Use System (SUS) Specialists



MSAT, VALIDATION & COMPLIANCE INTEGRATION

- Process Validation Engineers (IQ/OQ/PQ)
- GMP Compliance & Deviation Investigators
- QA/Compliance Integration Engineers
- Manufacturing & QMS Systems Analysts



CROSS-FUNCTIONAL & TECH TRANSFER ROLES

- Technology Transfer & Scale-Up Engineers
- Production & Process Development (PD)
 Experts
- Materials Management & Logistics Coordinators
- Manufacturing Training & SOP Development Specialists

Systems & Technologies That Drive Manufacturing Success

- Manufacturing Execution & Automation: Werum PAS-X, SAP, Oracle, DeltaV, Rockwell, Siemens PCS
- Quality & Compliance Platforms: TrackWise, MasterControl, Veeva Vault
- Lab & Analytical Tools: LIMS, Empower, ChemStation

Dexian's Purpose-Built Delivery Process















1. Discovery & Role Alignment

Define operational and compliance needs tied to scaleup, aseptic process control, and cGMP readiness.

2. Targeted Talent Sourcing

Identify candidates with handson GMP experience using Aldriven tools and deep industry networks.

3. Technical & Regulatory Verification

Screen for domain knowledge, certifications, and track records in pharma-grade manufacturing.

4. Seamless Integration

Support onboarding, documentation, and compliance processes for a smooth path to productivity.

Why Dexian?

SCALE WITH CONFIDENCE

Build high-functioning manufacturing teams—from clinical trial supply to commercial readiness.

MULTI-DISCIPLINARY ALIGNMENT

Talent with cross-functional fluency in Quality, Tech Transfer, and Supply Chain collaboration.

REGULATORY READINESS

Resources trained to meet FDA, EMA, and global pharmaceutical production standards.

CASE STUDY:

Scaling a 25+ Person Manufacturing Suite

CHALLENGE

A top pharmaceutical client needed to ramp up GMP manufacturing to meet commercialization demands.

SOLUTION

Dexian placed 25+ professionals, including aseptic operators, cell therapy technicians, and process engineers—while ensuring regulatory alignment and functional collaboration.

RESULT

Production scaled on time, with full compliance and operational success.

Let's build your GMP-compliant manufacturing team, together.