ISO 9001 Group

Consulting with a Personal Touch

Good Manufacturing Practices (GMPs)

Contact Us Today to Achieve GMP Compliance

The ISO 9001 Group understands the U.S. Food and Drug Administration's (FDA) Good Manufacturing Practice (GMP) regulations and how to comply with them. Our consultants can assist manufacturers, processors, and packagers of pharmaceuticals, medical devices, food and cosmetics ensure the safety, quality, and efficacy of their consumable products.

BENEFITS OF GOOD MANUFACTURING PRACTICE (GMP)

- ✓ Ensure consumer safety
- ✓ Comply with federal and statutory laws and regulations
- ✓ Ensure product safety, quality and efficacy
- ✓ Reduce consumer risks
- ✓ Meet requirements for GMP compliance
- ✓ Improved structure, processes and procedures internally

CLIENT TESTIMONIAL

"The responsiveness and communication provided by The ISO 9001 Group throughout our project was great. We liked the their approach on the technical support and guidance that was provided during the implementation sessions. Overall, the project went well and the meetings with our consultant were very helpful."

Ramesh Gopalakrishna
Saint-Gobain

HIGHLIGHTS OF 21 CFR PART 211

- ✓ Subpart A – General provisions
- ✓ Subpart B – Organization and Personnel
- ✓ Subpart C – Buildings and Facilities
- ✓ Subpart D – Equipment
- ✓ Subpart E – Control of Components and Drug Product Containers and Closures
- ✓ Subpart F – Production and Process Controls
- ✓ Subpart G – Packaging and Labeling Control
- ✓ Subpart H – Holding and Distribution
- ✓ Subpart J – Records and Reports
- ✓ Subpart K – Returned and Salvaged Drug Products

CONNECT WITH US

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