Providing Sterilization & Laboratory Services for the world's most innovative healthcare companies.

Announcement

Medistri Officially Listed Within the Swiss GLP Compliance Monitoring Program

May 18th 2026

Medistri is proud to announce that our laboratory is now officially listed within the Swiss GLP Compliance Monitoring Program under the oversight of the Swiss Federal Authorities and Swissmedic.

Following a successful inspection process conducted under the Swiss GLP framework, Medistri is now part of the national program responsible for monitoring laboratories operating in accordance with OECD Good Laboratory Practice (GLP) principles.

A Unique Position Within the Swiss GLP Landscape
The Swiss GLP Compliance Monitoring Programme includes a limited number of facilities operating under GLP oversight across Switzerland. Most facilities within the programme specialize in areas such as pharmaceutical toxicology, ecotoxicology, pathology, or bioanalytical chemistry.

Within this landscape, Medistri occupies a unique position as the only GLP laboratory in Switzerland combining:

- Chemical Testing
- Cytotoxicity
- Microbiology
- Cleanliness Testing

Within a single GLP-compliant laboratory infrastructure dedicated to the medical device and pharmaceutical industries.

This integrated approach allows manufacturers to consolidate multiple safety-critical testing activities with a single laboratory partner operating under both ISO 17025 accreditation and GLP compliance.

Strengthening Medistri's Laboratory Infrastructure
Medistri's official inclusion within the Swiss GLP Compliance Monitoring Programme further strengthens our laboratory operations and reinforces our commitment to supporting medical device, pharmaceutical, and biotech manufacturers with high-quality testing services operating under internationally recognized compliance frameworks.

Combined with our existing ISO 17025 accreditation, this milestone reflects the continued development of Medistri's laboratory infrastructure in Switzerland and our long-term commitment to scientific rigor, operational consistency, and regulatory reliability.

Our official GLP Certificate is now also available on our website.

To learn more about Medistri's laboratory services and GLP scope, please contact our dedicated team at lab@medistri.com or visit the official Swiss GLP register maintained by the Swiss Federal Authorities.

— The Medistri Team

#Medistri

This official registration confirms Medistri's position as the only GLP laboratory in Switzerland combining multiple safety-critical testing activities within a single GLP scope dedicated to the medical device and pharmaceutical industries. The official list of GLP-compliant laboratories in Switzerland, published by the Swiss Federal Authorities and now including Medistri as part of the Swiss GLP Compliance Monitoring Program, can be consulted here.

What This Means for Manufacturers
As regulatory expectations continue to increase across global healthcare markets, manufacturers are facing growing pressure to generate reliable, traceable, and internationally accepted safety data while maintaining efficient project timelines. By combining chemical testing, cytotoxicity, microbiology, and cleanliness testing within one GLP laboratory infrastructure, Medistri helps simplify the management of complex testing programmes. For manufacturers, this contributes to:

- Reduced coordination between multiple laboratories
- Greater consistency across testing activities and documentation
- Improved traceability throughout studies
- More streamlined project management
- Enhanced confidence during regulatory submissions

Studies performed under GLP conditions follow internationally recognized quality principles designed to ensure data integrity, reproducibility, and controlled documentation throughout the entire testing process.