TEMIS

Digital document management & documentation control system

TEMIS is a comprehensive Excel-Addin for the controlled preparation and administration of digital (GMP) documents

simple

Based on MS Excel architecture, no extra software, extensive manual provided

compliant

Validated according to FDA and EMA requirements for digital documentation

affordable

One licence per company, quick implementation, next to no training required

Why TEMIS?

Many small- and medium-sized pharma companies struggle with the demanding requirements for dealing with digital documents.

Digital documentation systems quickly exceed six digits in ($ or €), which is to many too big of an investment. Companies are either forced to fall back on paper-based documentation or resort to a cheap system (or none at all), which eventually does not comply with requirements.

This is where TEMIS comes into play. TEMIS allows document management according to GMP workflows, complies with the requirements, is validated, and does all that in just MS Excel and a Windows environment.

About QBDC

QBDC GmbH (Quality Biotech Development & Cells) is an international, Switzerland-based consulting company.

We are experts on the quality, development, and production of biosimilars, NBEs, and ATMPs. We facilitate your passage across technological, regulatory, and cultural gaps and are an aide in building a strong and sturdy GMP foundation.

With our vast experience in biotech topics, we can help you recognize risks before they become serious, help you prepare for challenges, help you optimize your workflows.

Eventually, your goal is to sell a authority compliant product; we can get you there, and get you there sooner.

visit qbdc.ch for more information

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