

**Proteos Biotech achieves
audit readiness and
enhanced compliance
with Scilife's eQMS**

Scilife



**See the
Customer Story**

About Proteos Biotech

Proteos Biotech, founded in 2010, specializes in recombinant enzymes and CMO services for biomedicine, pharmaceuticals, and cosmetics. The company is known for its innovative, high-quality products and commitment to regulatory compliance, partnering with clients to deliver tailored solutions across its diverse offerings.



Industry

BIOTECH

Regulations / Standards

GMP

Uses Scilife for

DOCUMENT CONTROL

TRAINING MANAGEMENT

EVENTS

CAPA MANAGEMENT

CHANGE CONTROL
MANAGEMENT

KPI

The Challenge

The company faced significant challenges in maintaining compliance and managing its growing document base using a paper-based quality management system. As they expanded rapidly, the limitations of manual processes became increasingly evident. To maintain compliance and ensure audit readiness, Proteos Biotech realized the need for a more efficient, scalable digital solution to streamline its quality management processes.

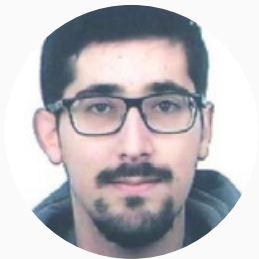
The Solution

To address these challenges, Proteos Biotech transitioned to Scilife's Quality Management System (QMS), which allowed them to centralize their document control, CAPA (Corrective and Preventive Action) management, and training processes. Scilife provided an integrated platform that met all their quality management needs, ensuring their operations remained compliant with industry standards.

Thanks to the dedicated support from Scilife's customer success team, the implementation of the QMS was both quick and smooth. Scilife's intuitive design and comprehensive functionality allowed Proteos Biotech to digitize its previously manual processes with minimal disruption to its daily operations.

“Before using Scilife, we stored all our documents in paper archives. Retrieving specific SOPs during audits was time-consuming, and we were constantly running out of storage space.”

Jorge Caño Martinez,
Quality Assurance
Technician.



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Quality Assurance
Technician.



“Now, during audits, we can access documents within seconds. This has reduced audit preparation time significantly.”

The Results

“Scilife has made compliance much easier. During audits, external auditors are always satisfied with how quickly we can present validated documentation.”

Maria Vendrell Moreno,
QA Manager.



Efficient Document Management

With Scilife's centralized platform, Proteos Biotech now efficiently manages its SOPs, CAPAs, and other critical documents. This has dramatically reduced the time spent searching for documents and preparing for audits. The system's robust search functionality and automated version control further ensure that the most current documents are always accessible, minimizing disruptions in daily operations.



Enhanced Compliance

The digital system has streamlined Proteos Biotech's ability to comply with industry regulations. Scilife's audit trail feature ensures full traceability of document changes and approvals, while its document validation capabilities keep Proteos Biotech prepared for any external audit. The comprehensive compliance reporting tools also facilitate easier preparation for regulatory reviews and inspections.



Improved Cross-Site Collaboration:

Scilife has improved communication and collaboration across Proteos Biotech's multiple sites in Spain by providing centralized access to all quality-related information. Teams now share and access critical data in real time, reducing errors and improving workflow efficiency. The system's integrated training management module also ensures consistent knowledge and procedures across all locations.

Reach Smart Quality with Scilife

Ready to bring new clarity to your Quality Management?

We're here, ready to help you turn quality into your brightest asset. Join esteemed leaders in the life sciences industry, such as Novartis, Biocartis, Yusen Logistics, Polpharma, Pendulum, and many more who trust Scilife to automate their processes, empower employees to take ownership of quality, enhance efficiency and transparency, and reduce compliance risk. The impact? Focus on what truly matters—creating transformative, life-saving solutions.

We'd love to hear from you!

Get in touch

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