MANAGEMENT OF THE LABORATORY EQUIPMENT MOVING PROCESS

An Application to manage the move of equipment from several laboratories to a single new building.

CHALLENGE

Identify all the steps required to move equipment, and structure an associated project management approach. Through various and powerful indicators, the solution must allow to easily manage the different activities, and the traceability of the equipment with labels integrating an ID and the information of origin and destination.

SOLUTION

Implementation of the GxpXL application to build a mock-up of the application.

Key users were able to align the vision of their needs with the possibilities of the GxpXL platform, then test and evolve the application according to feedback.

The application allows the equipment to be grouped into batches and to be tracked and monitored from its start location to its end location.

BENEFITS

The application allows the client to :

- Gain in productivity
- Optimize the monitoring of equipment and batches
- Facilitate the entry and research of information concerning a batch or equipment
- Trace all modifications/deletions/creations of information thanks to the audit trail
- Extract data in standardized and controlled formats
- Manage mandatory information entries
- Exploit the data entered to edit reports in Word or PDF and Excel format, depending on the rights
- Secure access to the application and its content
- Have an outline and indicators of the activity
- Have an overview of the moving activities

IN A FEW WORDS

CHALLENGES

- Easily manage the different activities
- Facilitate equipment traceability
- Data integrity

BENEFITS

- Optimization of equipment and batch monitoring
- Use of the audit trail
- Overview of activities
- Data security
- Performance Indicator

GxpXL

The GxpXL custom application allows you to protect your data and ensures compliance and agility for your projects by transposing classic files (Excel, Word, Lotus Notes, Visual Basic©…).

COMPLIANCE

- Compliance 21 CFR Part 11
- To international regulations : EMA, FDA, GMP, ISO
- Qualified application

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