



The industrial solution for manufacturers in the pharma- ceutical industry

*casyp*harm

The ERP solution for the pharmaceutical industry is called casyPharm. This software package is a validable solution that can be validated according to the requirements of GAMP-5, GxP, and FDA21 CFR Part 11. We can offer efficient and cost-effective solutions for performing system validation.

Experience is our competence **casymir.com**

CASYMIR

With our long industry experience, we have developed CASYPHARM, an ERP software that is tailored to the requirements of companies in the pharmaceutical industry. CASYPHARM offers a transparent representation of the entire process chain in all operational areas and is consequently batch-oriented. Certifications according to GxP, GAMP V, IFRA, REACH, VOC, FDA and other national regulations are possible and have been already implemented by our customers.

GMP compliant and validable

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HIGHLIGHTS

- Validatable system
- Update procedure to maintain validation
- Stability database to monitor product stability
- Logging of the production processes
- Active substance and substance management



RELEVANT FUNCTIONS

- Alidatable system
- Traceability of all data and processes
- Recipe management
- Flexible creation and management of certificates
- Ingredients, properties, areas of application of articles
- Acquisition and management of measurement data
- Easy management and documentation of free goods
- Flexible control of production facilities
- Ingredient management
- Support for order and stock production
- Calculation of product properties
- Integrated LIMS functions
- Stability database for product stability monitoring
- Secure product related warehouse management
- Maintenance of production facilities
- Multilevel production and release procedures
- Deposit of specifications
- Continuous traceability in all relevant areas
- Consistent workflow management
- Country- and customer-specific labeling with barcode GS1-128 (EAN)
- Batch traceability and quality management
- Customs Interface for Import and Export (CH)
- Complaints management
- Depiction of by-products
- Integrated weighing system
- Revision and release security
- Integration of document management systems
- Consignment stock and foreign production
- Audit Trail

CASYMIR Team Switzerland

 +41 61 716 92 22
 casymir schweiz ag
Fabrikmattenweg 11, CH-4144 Arlesheim

CASYMIR Team Germany

 +49 6834 9217-0
 SIGMATECH Informatik GmbH
Saarbrücker Str. 69, D-66359 Bous