SYSPRO for Pharmaceuticals

Research & Development

- Drug discovery
- Drug formulation
- Formulation adjustments
- Application for classification and registration
- Clinical trials
- Regulatory agency approval
- Patent registration

Drug approval and registration

Ingredient Sourcing

- Ingredient and material sourcing
- Ingredient traceability and tracking
- Receiving and inspections
- Compliance to regulatory warehousing storage
- Controlled area access

Audited issue of controlled ingredients

Product Manufacture

- Batch job creation
- Traceability and batch control
- Quality systems regulation
- Testing and inspection
- Allocation of lots and serial numbers

Warehousing

- Two-stage receipt and inspection
- Warehouse management
- Compliance to regulatory warehousing storage
- Control of lots and serial numbers
- Controlled area access

SOP

- Order taking via representatives
- Order management and releases
- Documentation with lots and expiry dates
- Audited order despatch
- Authorised P.O.D.

Customer Management

- Returns and recalls
- Sales calls and product education
- Analysis of customer buying trends
- Service and repair

Market Research

Warehouse Put-Away

Forecasting and planning

SYSPRO integration Framework
Automated feedback into Regulatory Requirement Applications

Administration
General Ledger, Cash Book, EFT, Assets, System Administration, SYSPRO Reporting Services

Bill of Materials
Engineering Change Control

Purchase Orders
Requisitions
Lot Traceability
Serial Tracking
Inspections
Warehouse Management
Quality Management

Work in Progress
Lot Traceability
Serial Tracking
Inspections
Quality Management

Quality Management
Warehouse Management
Inventory Control

Forecasting
Material Requirements Planning

Sales Order Processing
POD
Despatch

RMA
CMS

SYSPRO™

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