



SOLMAN AND FOCUSED BUILD AT RECIPHARM

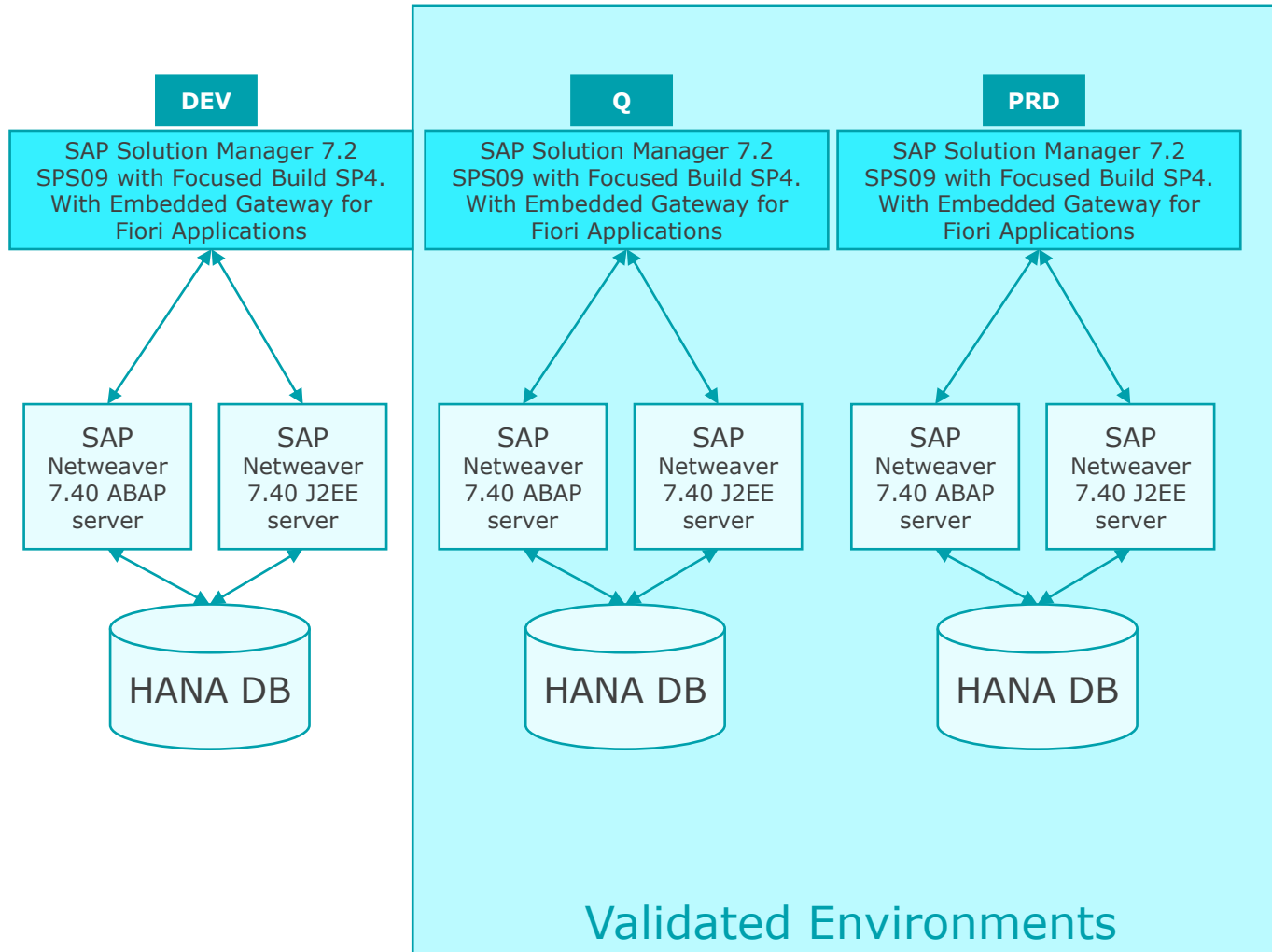
Patrick Cassini and Viktoria Engvall, Recipharm – 17SEP2021

AGENDA

- Scope of Solman in Recipharm
- Recipharm journey to Solution Manager
- Lessons learned



SCOPE OF SOLMAN IN RECIPHARM



3-tier landscape; Q and PRD validated

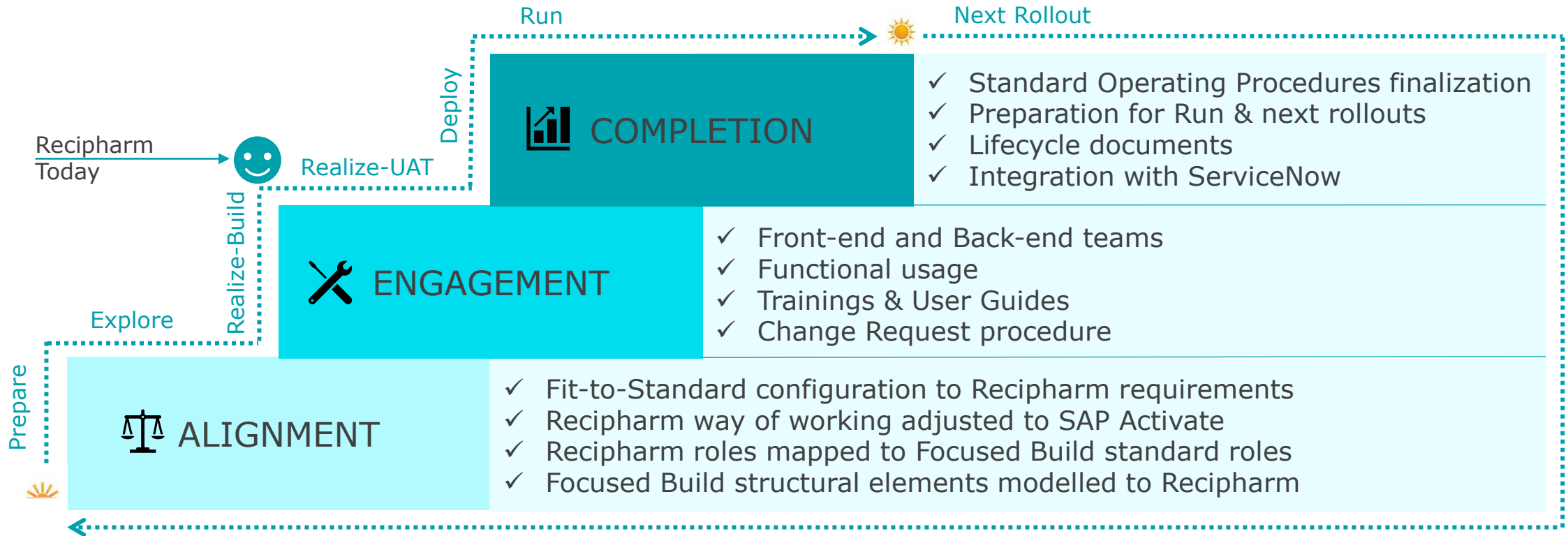
Focused Build – Requirements to Deploy

- To be used in the journey to S/4
- Solution Documentation
- Document repository for validation documentation
 - 2 level of signatures (business and QA) with strict authorizations
- Test Management for UAT (OQ/PQ)
- Release and change control management (To Be)
 - Fix Pace and Innovate

SNOW SAP Optimizer

- License optimization
- Connected to all ERP systems in the Group

RECIPHARM JOURNEY TO SOLUTION MANAGER

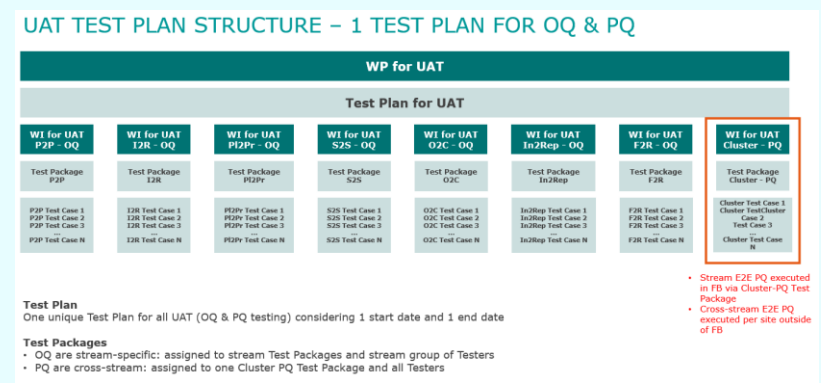


ALIGNMENT

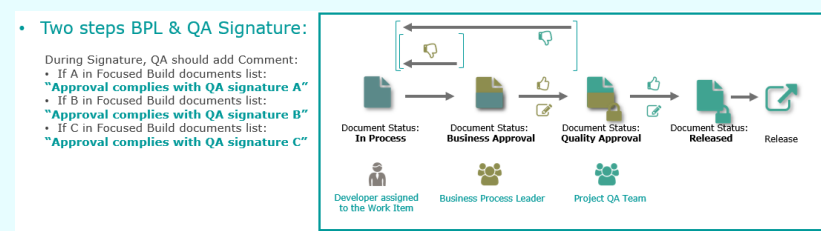


	FOCUSED BUILD	RECIPHARM
Way of Working	<ul style="list-style-type: none"> SAP Activate methodology 	<ul style="list-style-type: none"> When/who/where to upload and sign deliverables in Solman Definitions of FIT/GAP/WRICEF; Requirement/Defect classifications Approval procedures Quality Gates for phase ending: deliverables and status in Focused Build
Structural elements	<ul style="list-style-type: none"> Prepare: Solution-Project-Release Explore: L1-L5 SAP BP processes imported in Design Branch Realize-Build: Work Package-Work Item- Process-Transport Request-Documents Realize-UAT: Test Plan-Test Package-Test Case 	<ul style="list-style-type: none"> 1 Global Solution with Pilot Project and geographical rollouts L1-L4 in scope process selection and additional Interfaces & Master Data processes in Development Branch 1 FIT Work Package per Stream for standard customizing; 1 Work Package per WRICEF; L4 process and document assignment on Work Item level 1 overall Test Plan for UAT; 1 Test Package per Stream for functional testing; 1 Test Package for integration testing
Roles	<ul style="list-style-type: none"> Standard Roles 	<ul style="list-style-type: none"> Project Roles mapped to Standard Roles including internal Business Process Leads, Local Site Coordinators and Validation team members
Configuration	<ul style="list-style-type: none"> Fit-to-Standard (no development) 	<ul style="list-style-type: none"> Validation team defined as Role & Organizational Unit Validation repository for Validated documents 1-2 steps signature depending on Document Type Automated mail notification during UAT Recipharm Categories for Defects

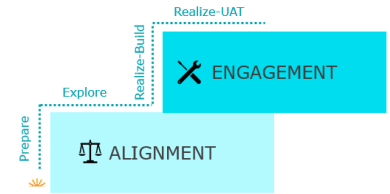
Document Type	Document Name	Process assignment in Focused Build	WP assignment	WP assignment	Document Release/Approval	QA	Document Approval/Release
ZPDS	Process Descriptions	BPH L4	N/A	N/A	N/A	NO	N/A
ZCSD	Conceptual Design	X.X.X.B "Stream documentation" L4 Process	Stream specific FIT Work Package "SEGR_XFX_B1"	Stream specific FIT Work Item "SEGR_XFX_B1"	Consultants	YES	N/A
ZTRC	Functional Design (L1 - S,C,E,F,W)	BPH L4	WRICEF WP	WRICEF WI	Consultants	YES	A
ZTRD	Technical Design (L1 - S,C,E,F,W)	BPH L4	WRICEF WP	WRICEF WI	Consultants	YES	A
ZQVD	Code Review	1. Recipharm Validation Documentation (non-process) repository L.X.X Validation Process	WP 2000000000 "Validation Documentation (non-process)"	WI 8000000010 "Validation Documentation (non-process)"	Consultants	YES	A
ZCSD	Configuration Specification	X.X.X.B "Stream documentation" L4 Process	Stream specific FIT Work Package "SEGR_XFX_B1"	Stream specific FIT Work Item "SEGR_XFX_B1"	Consultants	YES	N/A
ZQVD	Culture Plan Documents	1. Recipharm Validation Documentation (non-process) repository L.X.X Validation Process	WP 2000000000 "Validation Documentation (non-process)"	WI 8000000010 "Validation Documentation (non-process)"	Recipharm	YES	A
ZQVD	General Plan used in validation	1. Recipharm Validation Documentation (non-process) repository L.X.X Validation Process	WP 2000000000 "Validation Documentation (non-process)"	WI 8000000010 "Validation Documentation (non-process)"	Recipharm	YES	A
ZQVD	Data Migration Plan	1. Recipharm Validation Documentation (non-process) repository L.X.X Validation Process	WP 2000000000 "Validation Documentation (non-process)"	WI 8000000010 "Validation Documentation (non-process)"	Recipharm	YES	A



Recipharm Role \ Standard Roles	Business Analyst	Architect	Developer	Project Manager	Release Manager	Test Manager	Tester	Business signature	QA signature
Business Process Lead (BPL)			X					X	
Intelligence Team Lead	X		X						
Solution Architect		X							
Project Manager				X			X		
Local Site Project Coordinator (LSC)	X		X					X	
Key-User	X								
Project QA	X		X				X		X
Methodology Coach									
SAP CoE	X	X	X	X	X		X	X	
Licence Manager									



ENGAGEMENT



Back-End

Licence Manager

- Optimizes licenses for ERP

Service Owner

- Accountable for Focused Build

Service Manager

- Provides end-user trainings and user guides
- Establishes Standards of Procedures
- Tests system changes in Quality system
- Administrates user roles and authorizations



Solution Experts

- Provide continuous support including end-user training and user guides
- Set up Recipharm solution
- Implement SAP notes and Change Requests in Development system

PMO

- Use management operations
- Control status on progress and deliverables
- Align with business streams

Validation Team

- Sign documents for quality review
- Sign change requests
- Use validation repository

Requirements

Configuration
SAP notes
System upgrades

Recipharm Focused Build Application

Production

Front-End

Finance to Report

Inspect to Release

Order to Cash

Plant to Produce

Purchase to Pay

Store to Ship

Inspect to Repair

- Use Focused Build for business operations (scoping, development, testing)
- Upload and sign deliverables for business review

SAP CoE

- Use Focused Build for Change Management and Fix Pace post go-live
- Responsible for lifecycle documents

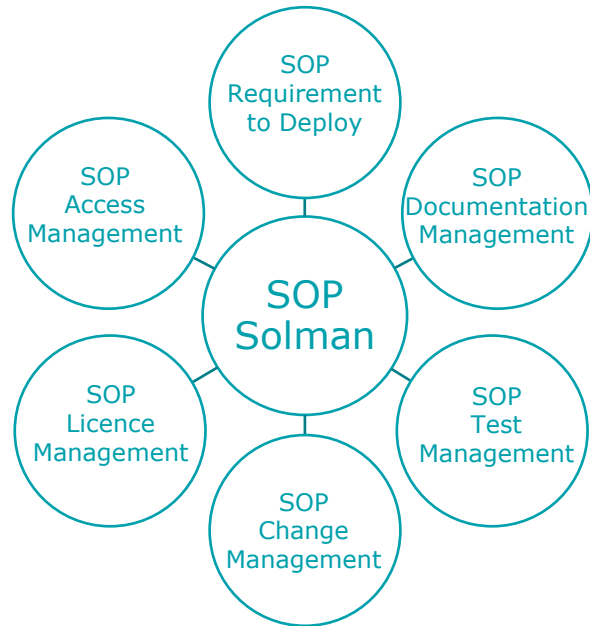
Internal team External team

COMPLETION



Standard Operating Procedures

Internal documents describe our ways of working, use and maintenance of our central SAP Solman system.

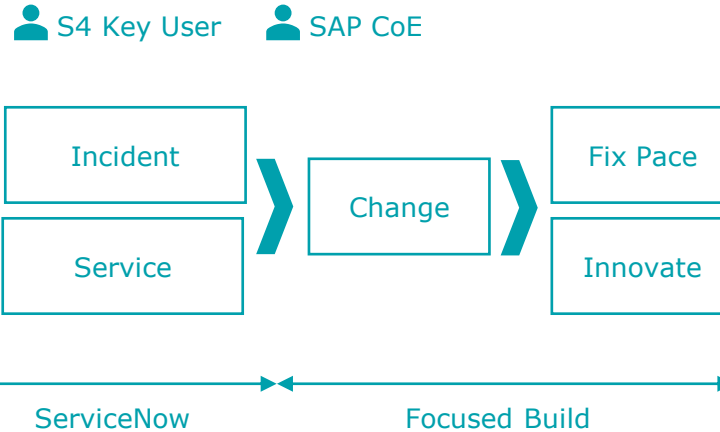


- ❖ Scope
- ❖ Roles and RACI matrix
- ❖ Definitions
- ❖ Process workflows
- ❖ Process descriptions
- ❖ User instructions



Run preparation

When S4 is going live, SAP CoE must be prepared to utilize Focused Build for short and long term changes.

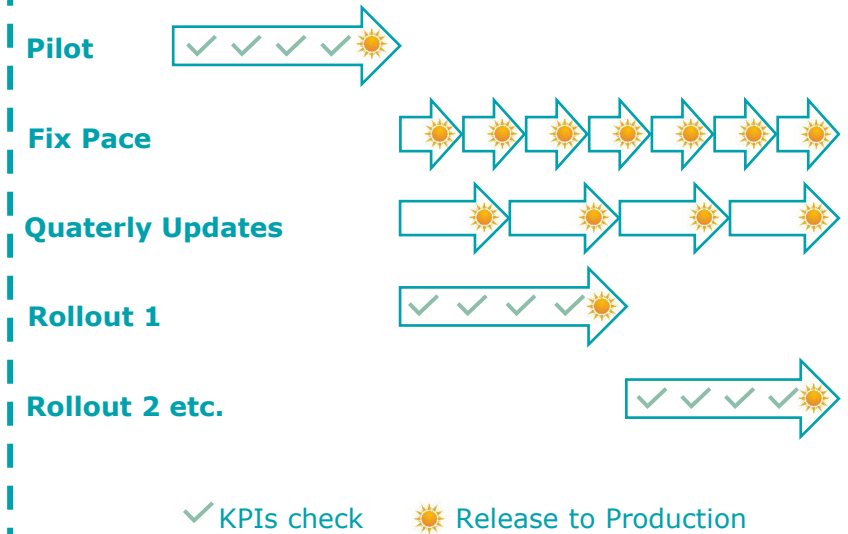


- ❖ Incidents & Services start from ServiceNow
- ❖ Fix Pace for quick fixes
- ❖ Business Requirements
- ❖ System changes start from Focused Build
- ❖ Innovate for release changes
- ❖ Lifecycle documentation

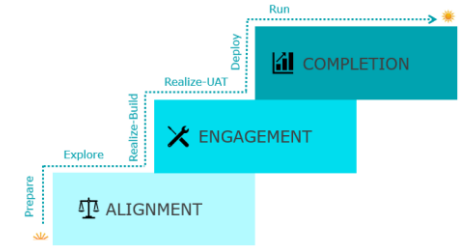


Next Rollouts preparation

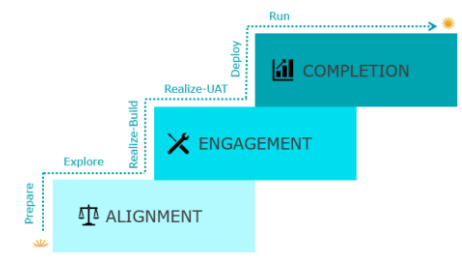
Lessons learned from first deployment will be used to get resources and system organized for next rollouts.



- ❖ Release-Project structure
- ❖ KPIs set up for deliverables
- ❖ Use of Requirements in Focused Build
- ❖ Optimized way of working
- ❖ Early training of internal/external teams
- ❖ Full involvement of the teams



EXAMPLE OF SOP - UAT PREPARATION

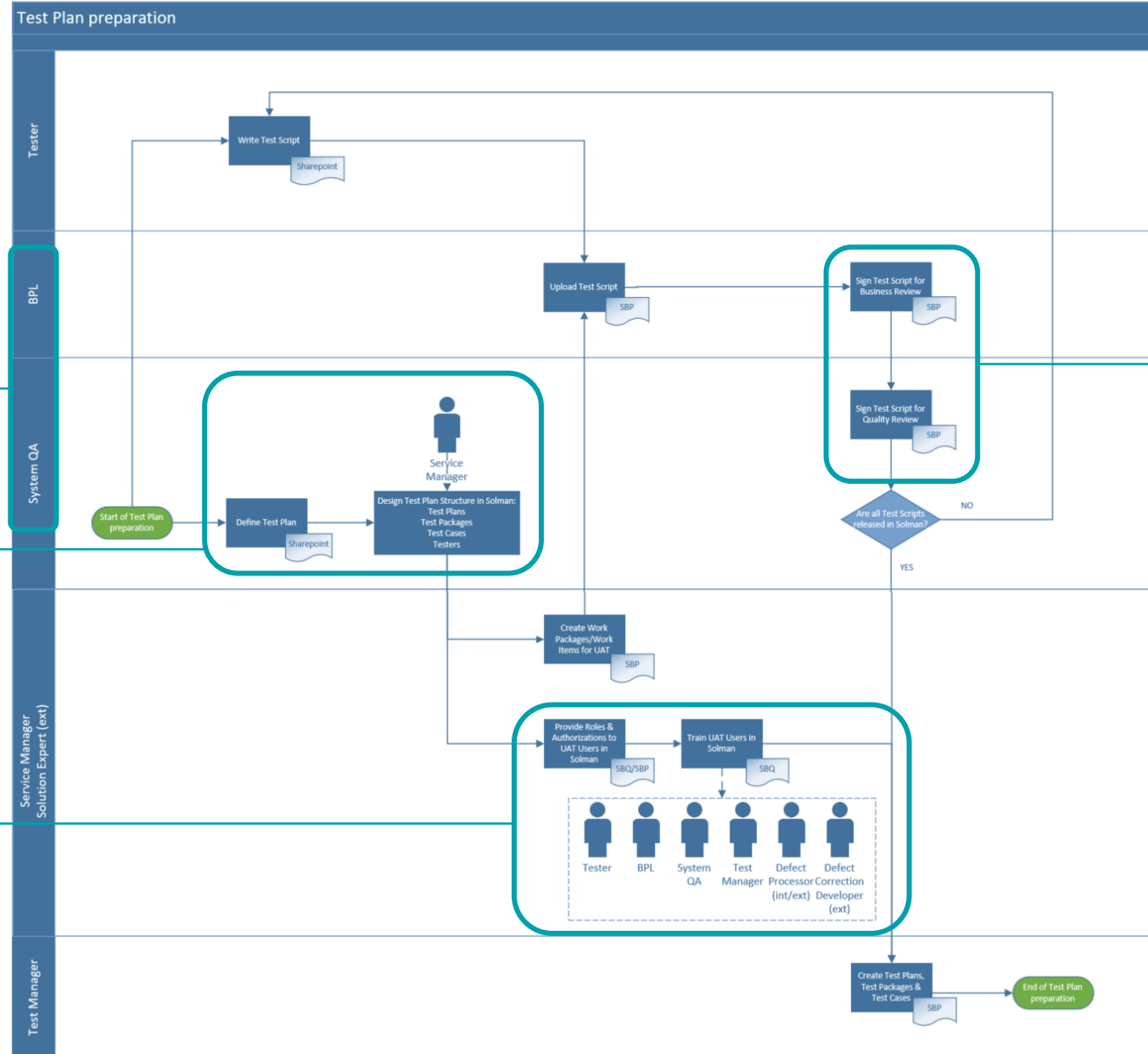


✓ Validation team & Business Process Lead roles

✓ Inclusion of functional & integration test plan following lifecycle validation risk-based approach

✓ Internal preparation

✓ Double step signature



LESSONS LEARNED

Key Challenges

- Lifescience industry requirements
- One global solution
- Internal resources & knowledge
- Documentation on L4 process level
- Use of Requirements
- SAP/lifescience Best Practices processes
- Missing assignments & deliverables
- Integration to ServiceNow

Key Opportunities

- Flexibility to fit our requirements
- One ALM tool for One global S4 solution
- Internal teams, SOP and User Guides
- Lifecycle documents on L4 level
- Recipharm own Way of Working
- Template solution
- Streams engagement and use of KPIs
- From ServiceNow to Focused Build

