****<organisation> Non-SIT S&A MHHS QT Test Plan****

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# Contents

[1 Contents 1](#_Toc198565367)

[1.1 Programme Participant Name 4](#_Toc198565368)

[1.2 Change Record 4](#_Toc198565369)

[1.3 Document Approval 4](#_Toc198565370)

[1.4 References 4](#_Toc198565371)

[1.5 Terminology 4](#_Toc198565372)

[2 Scope 5](#_Toc198565373)

[2.1 Scope Overview 5](#_Toc198565374)

[2.1.1 In Scope 5](#_Toc198565375)

[2.1.2 Out of Scope 5](#_Toc198565376)

[3 Test Approach 6](#_Toc198565377)

[3.1 Placing Reliance 6](#_Toc198565378)

[3.2 Test Scenarios and Cases 6](#_Toc198565379)

[3.3 Re-Testing and Regression 6](#_Toc198565380)

[3.4 Test Entry Criteria 6](#_Toc198565381)

[3.5 Test Exit Criteria 8](#_Toc198565382)

[4 Test Schedule and Deliverables 8](#_Toc198565383)

[4.1 Test Schedule 8](#_Toc198565384)

[4.2 Qualification Test Execution Plan (\*NEW\*) 9](#_Toc198565385)

[5 Test Infrastructure 9](#_Toc198565386)

[5.1 Test Environments 9](#_Toc198565387)

[5.2 Test Stubs and Other Tools 9](#_Toc198565388)

[5.3 Test Data 9](#_Toc198565389)

[5.4 Release and Configuration Management 9](#_Toc198565390)

[6 Roles and Responsibilities 9](#_Toc198565391)

[6.1 Test Roles and Responsibilities 9](#_Toc198565392)

[7 Defect Management 11](#_Toc198565393)

[7.1 Defect Management and Reporting of Non-SIT S&A Defects 11](#_Toc198565394)

[7.2 Work-Off Plan 11](#_Toc198565395)

[8 Test Result Management and Reporting 11](#_Toc198565396)

[9 Test Assurance 11](#_Toc198565397)

[10 Risks, Assumptions, Issues and Dependencies 12](#_Toc198565398)

[10.1 Risks 12](#_Toc198565399)

[10.2 Assumptions 12](#_Toc198565400)

[10.3 Issues 12](#_Toc198565401)

[10.4 Dependencies 12](#_Toc198565402)

[Appendix 13](#_Toc198565403)

**Guidance notes:**

Use of this template for documenting a QT Test Plan is recommended. A Programme participant may use another format if required but must ensure the same topics are covered in accordance to the guidance provided for each section of this document. Programme participants can also add any additional sections as they feel appropriate.

Orange text indicates guidelines or examples.

Template starts from page 4 onwards - this page must be removed from the final document produced by Programme participants

## Programme Participant Name

*Programme participants to fill in the table below and include their relevant constituency group and third-party provider if applicable.*

|  |  |  |  |
| --- | --- | --- | --- |
| Programme Participant Name | Constituency Group | MPID(s) Tested | Third Party Provider |
| *e.g. Generic Electricity Co. Ltd* | *Non-SIT S&A* | *MPID1* | *N/A* |

## Change Record

*The table below needs to be updated with details of each amendment and change control.*

|  |  |  |  |
| --- | --- | --- | --- |
| Date | Author | Version | Change Detail |
|  |  |  |  |
|  |  |  |  |
|  |  |  |  |
|  |  |  |  |

## Document Approval

*The table below needs to be updated and the role needs to be defined per person as one of the following categories:*

* *Review;*
* *For Information;*
* *For Approval; and*
* *Approval and Sign Off.*

|  |  |
| --- | --- |
| Reviewer/Approver | Role |
|  |  |
|  |  |
|  |  |
|  |  |

## References

*Reference any documents into the table below along with reference and version number. Programme participants should insert documents wherever possible.*

|  |  |  |  |  |
| --- | --- | --- | --- | --- |
| Reference | Documents | Publisher | Version | Additional Information |
| *REF 001* | *Requirements to Test Traceability Matrix* |  |  |  |
| *REF 002* | *QT Pro Forma/Test Catalogue* |  |  |  |
| *REF 003* | *QT Test Readiness Report* |  |  |  |

## Terminology

*Reference any terminology requiring a description used in this document.*

|  |  |
| --- | --- |
| Term | Description |
|  |  |
|  |  |
|  |  |

*.*

# Scope

## Scope Overview

*This section is intended to define the high-level scope of Non-SIT S&A QT. It should list areas of the system or functionality being tested and should include a reference to functional, migration, non-functional, operational requirements in scope and design baseline.*

*An architecture diagram should be included which clearly indicates all systems being used as well as test stubs or other test software. A description of limitations of any test stubs or software should be included along with the way these limitations are being mitigated.*

*NOTE that this is an important section of the document for assurance purposes.*

### In Scope

*The Scope for Qualification Testing should be defined by your draft RTTM and Test Catalogue; please ensure these are provided with this document.*

### Out of Scope

*This section should explicitly identify the high-level key features, test requirements and interfaces that will not be tested by listing what is out of scope of the test plan along with reasons for not testing. This should also include any DIP publications that parties are opting out of.*

|  |  |  |  |
| --- | --- | --- | --- |
| Key Themes/Requirements/Interfaces | Test Types | Risk Level | Provide Rationale |
| Theme x |  |  |  |

Table 2 Out of Scope

# Test Approach

## Placing Reliance

*This section describes your placing reliance approach if applicable. This is expected to be a high-level summary unless there are any changes to the placing reliance approach already shared.*

*Provide your response below:*

## Test Scenarios and Cases

*As per Section 2.1.1, the RTTM should identify all test cases as identified as ‘in scope’ from the Test Catalogue, also ensuring that the Test Catalogue has been annotated to demonstrate any exclusions due to exemptions or placing reliance (please ensure this is also reflected in the RTTM). Any additional information or context to support the RTTM should be added below:*

## Re-Testing and Regression

*This section should describe a Non-SIT S&A approach to any retesting that will be expected to be carried out due to defect, clarification process etc., including risk assessments and regression testing against IR8.x where parties have been unable to complete against the current version deployed for QT as part of PIT. Please refer to the most current version of MHHS-DEL852 - Pre-Integration Testing Guidance for detailed advice on regression and risk assessment and explain how this will be managed prior to Qualification Testing start.*

## Test Entry Criteria

*This section describes how test entry criteria defined in Annex 2 Non-SIT S&A QT Approach and Plan section 10.6 will be met. Please describe how the following criteria will be carried out and evidenced (where applicable) prior to QT start:*

|  |  |
| --- | --- |
| **​** | **​Non-SIT S&A Entry Criteria** |
| ​1 | ​Evidence of successful PIT Completion has been submitted, assured by the Non-SIT S&A QT Team and any work off plans agreed with Code Bodies and tracked - please refer to [MHHS-DEL852 - Pre-Integration Test Guidance](https://mhhsprogramme.sharepoint.com/sites/Market-wideHalfHourlySettlement/Testing%20Documents/MHHS-DEL852%20-%20%20Pre-Integration%20Testing%20Guidance%20v1.1.pdf?web=1) for full details of the PIT exit criteria and submission timelines for PIT deliverables. |
| ​2 | ​Non-SIT S&A Participant are on track to submit evidence of successful PIT Completion (non-functional and operational) by the agreed dates as outline in the Programme Plan. |
| ​3 | ​All relevant test artefacts listed as an entry criteria in section 10.4 of Annex 2 (MHHS-DEL2433) must have been produced by the Non-SIT S&A QT participant and approved by Non-SIT S&A QT Team and Code Bodies (where applicable). |
| ​4 | ​Successful completion of DIP onboarding and connectivity proving with no open issues which would impact the test schedule for Non-SIT S&A QT. |
| 5 | Successful completion of onboarding into ADO with confirmation of resource access |
| 6 | Successful completion of onboarding into QTF with confirmation of resource access including completion of smoke test |
| ​ |  |
| ​7 | ​Test Data (Test MPANs) has been requested, generated and loaded into the Participant UIT environment. |
| 8 | ​Participants have confirmed they have resources with necessary skills and system access to support the test stage execution, the defect management process and have completed QTF/ADO training. |
| ​9 | At least one primary contact has been provided and onboarded into the MHHSP Microsoft Teams private Participant channel for the purpose of Qualification Testing. |
| ​10 | ​For any Non-SIT S&A QT participants placing reliance, MHHS Placing Reliance submission reviewed and approved by Non-SIT S&A QT Team and Code Bodies. |
| ​11 | ​Where test scope has been adjusted for Placing Reliance and/or exemptions have been requested, this has been documented in the Section 3.1 of this document and this has been agreed by the Non-SIT S&A QT Team and Code Bodies.  The Participant will be required to reflect test case exemptions in the RTTM and submitted Test Catalogue in accordance with the Placing Reliance agreement. |

## Test Exit Criteria

*This section should how test exit criteria defined in Annex 2 Non-SIT S&A QT Approach and Plan section 11.8 will be met.*

|  |  |
| --- | --- |
|  | *Exit Criteria* |
| *1* | No outstanding Severity 1 and Severity 2 Defects. |
| *2* | Sev3 and Sev4 defects that cannot be resolved during QT are documented with impacts assessment and a work off plan by the Programme Participant, reviewed by the Non-SITS&A QT Team and agreed with Code Bodies and DIP Manager ahead of QT exit. |
| *3* | 100% test execution coverage, including functional, non-functional, migration and operational testing with approval from Code Bodies and DIP Manager for any de-scoped/ failed test cases prior to QT completion. |
| *4* | Non-SIT S&A QT Completion Report including work off reviewed by Non-SIT S&A QT Team and submitted to Code Bodies and DIP Manager. |
| *5* | Test results and evidence have been uploaded to the test management tool (QTF) and has been assured by Non-SIT S&A QT Test Team and approved by Code Bodies. |
| *6* | Non-SIT S&A QT Completion Report, including work-off plans approved by Code Bodies and DIP Manager, reviewed and approved by the Non-SIT S&A QT Team and submitted to Code Bodies and DIP Manager. |

# Test Schedule and Deliverables

## Test Schedule

*This section should provide details of the proposed high-level test schedule regarding the testing activities. This information will be used by the Non-SIT S&A QT Team to track Participant progress. It should also include details of:*

* *Test preparation (including RTTM, Test Catalogue and other deliverables from section 10.4 of Annex 2 Non-SIT S&A QT Approach and Plan)*
* *Test execution dates*

*Programme participants can use the table below as an example or create a new table based on their test stage testing schedule:*

|  |  |  |
| --- | --- | --- |
| Test Activities | Start Date | End Date |
| *e.g., Produce RTTM* | *01/03/2023* | *01/04/2023* |
| *e.g., Provide Test Catalogue* | *01/03/2023* | *01/04/2023* |
| *Add additional lines as required….* |  |  |

Table 3 Test Schedule for test stages

## Qualification Test Execution Plan (\*NEW\*)

*Please complete the tables below to indicate how the qualification test set will be completed within the assigned test wave on a fortnightly basis. This will be monitored and reported by the Non-SIT S&A Team during Qualification Testing as per OFGEM requirements.*

|  |  |  |
| --- | --- | --- |
| Start Date | End Date | Total Test Cases |
| dd/mm/yyyy | dd/mm/yyyy | *XX* |

|  |  |  |  |  |  |  |  |  |
| --- | --- | --- | --- | --- | --- | --- | --- | --- |
| Week | 1,2 | 3,4 | 5,6 | 7,8 | 9,10 | 11,12 | 13,14 | 15,16 |
| Planned Test Cases | *XX* | *XX* | *XX* | *XX* | *XX* | *XX* | *XX* | *XX* |

# Test Infrastructure

*This section should define the environment to support the preparation and execution of this Non-SIT S&A QT Test Plan. Participants should also identify systems where Placing Reliance has been applied e.g. DIP adapter, UMS exemptions.*

## Test Environments

*This section should define the Programme Participant's test environments used within QT stage.*

## Test Stubs and Other Tools

*This section should define any test stubs and other tools a Non-SIT S&A is using within QT stage that is not the QTF or MHHSP Test Management Tool (ADO).*

## Test Data

*This section should include any further information to Non-SIT S&A’s approach to test data that is not documented in the Non-SIT S&A QT Test Data Approach and Plan.*

## Release and Configuration Management

*This section should define the release and configuration management approach for a Non-SIT S&A’s systems to be used within QT stage. This should include details about how regression is evaluated and carried out.*

# Roles and Responsibilities

## Test Roles and Responsibilities

*In this section, define the key roles needed to fulfil QT testing. Clear and unambiguous definition of the roles and responsibilities is essential for fulfilling this Test Plan. If possible then Programme participants can put a name against each role as well as the job title. One person may hold more than one role* *providing there is no conflict of interest.*

*The table below* *identifies the roles and responsibilities for QT stage:*

|  |  |  |
| --- | --- | --- |
| Role | Full Name (optional) | Responsibilities |
| *e.g., Test Manager* |  | *Day to day test management and progress reporting, review of test inputs and test outputs, allocation of test tasks* |

Table 4 Test Roles and Responsibilities

*Test team structure of the test organisation can be added if available*.

# Defect Management

## Defect Management and Reporting of Non-SIT S&A Defects

*This section should describe the defect management process for defects raised by Non-SIT S&A Programme participants on their systems.*

## Work-Off Plan

*This section should describe the approach to using a work-off plan, should one be required.*

# Test Result Management and Reporting

*This section should describe how reporting to the Non-SIT S&A Team, DIP Manager and Code Bodies for QT readiness will be performed.*

# Test Assurance

*This section should describe the test assurance process in detail and how Programme participants will carry out self-assurance on all their test-related deliverables and activities for QT.*

*It should also describe any criteria for a work-off plan; a template is planned and will be distributed.*

# Risks, Assumptions, Issues and Dependencies

## Risks

*The focus within this section should be on risks that relate to and affect testing in this Test* *Plan and its scope. Any non-testing related risks* *identified by the Programme participant should also be listed below.*

|  |  |  |  |  |
| --- | --- | --- | --- | --- |
| Risk No | Risk Description | Impact | Owner | Mitigation |
| *R-1* | *Insert description* | *Describe how the risk could affect testing* | *Set an owner* | *Describe what steps are being taken to mitigate the risk* |

Table 5 Risks

## Assumptions

*The focus within this section should be on assumptions that relate to and affect testing in this QT Test Plan and its scope. Any non-testing related assumptions* *identified by the Programme participant should also be listed below.*

|  |  |  |
| --- | --- | --- |
| Assumption No | Assumption Description | Rationale |
| *A-1* | *Describe the assumption* | *Summarise why the assumption is needed* |

Table 6 Assumptions

## Issues

*The focus within the section should be on issues that relate to and affect testing in this QT Test Plan and its scope. Any non-testing related issues* *identified by the Programme participant should also be listed below.*

|  |  |
| --- | --- |
| Issue No | Issue Description |
| *I-1* | *Describe the issue* |

Table 7 Issues

## Dependencies

*The focus within this section should be on dependencies that relate to and affect testing in this QT Test Plan and its scope. Any non-testing related dependencies* *identified by the Programme participant should also be listed below.*

|  |  |
| --- | --- |
| Dependency No | Dependency Description |
| *D-1* | *Describe the dependency* |

Table 8 Dependencies

# Appendix

*Any appendix should be added here.*