

Quality assurance for Manufacturing

Dressed Cavities for SPS prototype cryomodule



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Project schedule

Presented during Crab Cavities Workshop, December 2013

	2014				2015				2016			2017				
	Q1	Q2	Q3	Q4	Q1	Q2	Q3	Q4	Q1	Q2	Q3	Q4	Q1	Q2	Q3	Q4
First cryomodule	Yellow	Orange	Orange	Orange	Light Green	Light Green	Pink	Purple	Blue	Light Blue	Light Blue	Green	Green			

↑ Today

Constraint: fixed time window for SPS installation

Cryomodule Design	Dark Red	Red	Olive													
Cleanroom Activity			Grey	Grey	Black											

Yellow	cavity engineering design and interfaces	Purple	cavities assembly in clean room
Orange	cavities technical design and manufacture	Blue	cryostating
Light Green	bare cavities processing and tests	Light Blue	cryomodule tests in SM18
Pink	helium tank assembly on cavity	Green	ready for SPS installation

Dark Red	cryomodule engineering design (incl helium tank+tuner)	Grey	clean room assembly and cryostating tooling design
Red	cryomodule technical design	Dark Grey	clean room tooling manufacturing
Olive	cryomodule manufacturing	Black	cryostating tooling

Requirements for Manufacturing Readiness

Due 1 month before start of manufacturing for CERN approval

#	Requirements	Y/N
1	Niobium material samples according to Section 3.2	N
2	Material certificates and quality control of raw materials (including RRR measurements)	N
3	Material certificates of welding consumables (whenever applicable)	N
4	Functional and manufacturing drawings (with tolerances)	Y?
5	Design reports demonstrating that welds are designed to withstand the specified load cases (refer to Section 3.6.1)	N
6	Welding plan including: <ul style="list-style-type: none"> • Welding maps • Welding and brazing procedure qualification record including CERN acceptance criteria in Section 4.2 (WPQR and BPQR) • Welding and brazing procedure specification (WPS and BPS) • Welders performance qualification (GTAW), Welding and Brazing Operators Performance Qualifications (electron-beam welding and vacuum brazing) – WPQ, WOPQ and BOPQ 	N
7	Manufacturing procedures (whenever required in Annex 6.3)	N
8	Test procedures (whenever required in Annex 6.3)	N
9	EB welded and vacuum brazed samples according to the requirements specified in Section 3.8.4	N
10	NDT personnel qualifications	N
11	Manufacturing and inspection plan (MIP) – list of all manufacturing and quality control operations.	N

Mandatory manufacturing and quality control steps

Step ID	Step name	CERN control
ACFCA-1-QA	Shaping and machining	N/A
ACFCA-2-QC	Dimensional control	HP
ACFCA-3-QA	Cleaning and chemical polishing	N/A
ACFCA-4-QA	Vacuum brazing	N/A
ACFCA-5-QC	Visual examination	HP
ACFCA-6-QC	Dimensional control	HP
ACFCA-7-QC	Helium leak tightness test	NP
ACFCA-8-QC	Ultrasonic examination	HP
ACFCA-9-QA	Chemical polishing	N/A
ACFCA-10-QA	Electron-beam (EB) welding	N/A
ACFCA-11-QC	Visual examination	HP
ACFCA-12-QC	Radiographic examination	HP
ACFCA-13-QC	Dimensional control	HP
ACFCA-14-QC	Resonator frequency check & tuning	HP
ACFCA-15-QC	Dimensional control	HP
ACFCA-16-QC	Helium leak tightness test	N/A
ACFCA-17-QA	Packaging & shipping	HP
ACFCA-18-QA	Bulk chemical polishing	N/A
ACFCA-19-QA	Heat treatment	N/A
ACFCA-20-QA	Light chemical polishing	N/A
ACFCA-21-QA	High pressure water rinse	N/A
ACFCA-22-QC	RF acceptance tests at cold temperature	HP



Input for
CERN

**Any further steps
in the MIP shall be
communicated to CERN !**



Required specific
manufacturing
and test procedures

Quality controls points

According to Section 1.3 of the engineering specification

- **Notification Point (NP):** CERN, or its authorized representative, is informed 5 working days in advance that a specific step has been completed and that the following step in the approved work-flow will be performed. A NP does not affect the work-flow. Work can continue without CERN, or its authorized representative, reply.
- **Hold Point (HP):** CERN, or its authorized representative, is informed that a specific step has been completed. The work-flow is stopped until CERN, or its authorized representative, provides a HP Clearance. The clearance is provided within 5 working days upon submission of the quality control documentation relative to the performed step. In case of clearance the work-flow can continue. In case of rejection, a recovery plan shall be discussed with CERN and submitted to CERN for final approval within 10 working days.

Management of non-conformities

Nonconformity Report	
IDENTIFICATION	
1. Originator's Name:	6. Date:
2. Contractor/Supplier:	7. Part description:
3. Contract No:	8. Qty:
4. Project Engineer:	9. Dwg No:
5. Quality Manager:	
10. Found during what activity:	
<input type="checkbox"/> Incoming inspection	<input type="checkbox"/> Final inspection
<input type="checkbox"/> In-process inspection	<input type="checkbox"/> Other:
11. Description of nonconformity (Use continuation page if necessary)	
12. Action taken to prevent misuse (use continuation page if necessary)	
IMPORTANCE	
13. <input type="checkbox"/> Non critical	<input type="checkbox"/> Critical
DISPOSITION	
14. <input type="checkbox"/> Use-as-is	<input type="checkbox"/> Repair
<input type="checkbox"/> Reject	<input type="checkbox"/> Rework
<input type="checkbox"/> Return to supplier	
Description of proposed action (use continuation page if necessary)	
CORRECTIVE/PREVENTIVE ACTION	
15. Description of proposed action (Use continuation page if necessary)	
APPROVAL OF NON CRITICAL NONCONFORMITIES	
16. Project Engineer:	Date:
APPROVAL OF CRITICAL NONCONFORMITIES	
17. Project Management:	Date:
CLOSURE OF THE NONCONFORMITY	
Planned actions have been completed and corrective/preventive actions have been initiated	
For non critical nonconformities For critical nonconformities	
18. Name: Quality Manager or Project Engineer	Name: Project Engineer
Date:	Date:

According to Section 1.3 of the engineering specification

- Non-conformities observed during the specified quality controls in the different processes shall be appropriately managed and handled by the supplier. All reports of non-conformities, including corrective and preventive measures, shall be submitted to CERN for approval before implementation.

Non-conform parts/materials shall be appropriately labelled as “non-conform” and removed from the manufacturing chain

Conclusions

- Where we are today?
 - Schedule: about 3 months delay in the manufacturing process
 - Requirements for manufacturing readiness not fulfilled → CERN clearance for manufacturing can not be provided.
- What is the action plan from the Supplier to recover from delays/provide missing information as soon as possible?
- Where can CERN provide assistance?