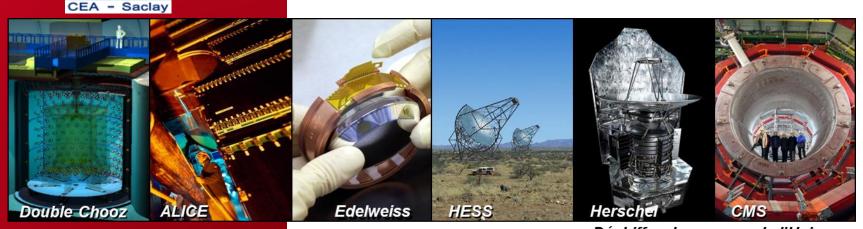
DE LA RECHERCHE À L'INDUSTRIE

PIP-II TECHNICAL WORKSHOP

CAVITY MANUFACTURING WG 01-04 DEC20

Topic 5- Quality Control of Cavity Production



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Vincent HENNION



Introduction

- Lessons learned gained from EXFEL and ESS Accelerator Facilities
- Production of high number of cryomodules for elliptical cavities
- Framework of quality control
 - different partners and contributors
 - responsibilities sharing
 - Construction phase (prototype vs series production)
 - Expertise level





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FRAMEWORK OF QUALITY CONTROL

Objectives

- Verification that specified requirements are fulfilled
- Ensure traceability of all deviations
- Contribute to configuration management

Consistency with system engineering

- Technical needs in different phases of the cavity lifecycle
 - Performances/specifications
 - Procedures/Adjustments
- Technical constraints
 - Processes/optimizations/risks
 - Intellectual properties

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TECHNICAL NOTE

Interfaces document High Beta cavities and cryomodules



Released by : Christophe NAYRI, Chef de prolet ESS-Date et signature :





Consistency with implemented organisation

- Prototyping phase
 - Technical challenges
 - Specification of the requirements for the series
 - Agreed way of working between partners
 - communication/information channels to be set up
- planning constraints
 - Critical path is not impacting all partners at the same time
 - The more partners and stakeholders there are, the more complex is the interfaces management
 - Higher is the number of interfaces, higher shall be the involvement and the effort by partners in Quality Control



Scope: elliptical cavity is a complex system

- Performances are high level requirements.
- Specs are defined from performances to reach.
- Optimization of the cavity cost is directly linked to the tolerances.
- Tolerances are directly linked to the manufacturing processes to be implemented.
- Processes to be implemented should be homogeneous (orderer shall discuss with the manufacturer as much as possible)



01-04/12/2020





Topic 5: Quality



How to successfully transfer and fulfill specs?

- Find technical solutions for mechanical assemblies
 - Relax some tolerances to extend acceptance capabilities
 - deviation impact analyses (dummy tools for tuner interfaces or cavity string interfaces with spaceframe)
- References transfer
 - Strong recommendation to specify references based on dimensions issued from milled parts (accuracy and reproducibility)
 - Impact on production cycle and process: technical risk
 management to be discussed with the manufacturer



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Thank you

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