RPC Purifiers' studies /status/

> IR, RG 11.01.2010

The task

Problems

- Possible high currents on long term operation of RPCs
- Small amounts of F based impurities observed
- Under irradiation larger amounts and additional impurities observed
- Constant water levels in the gas mixture vital for the RPC operation





The **Solution** we search is..

The optimal combination of filters:

- With the best H2O and O2 filtering capacity
- Removing maximum F-based impurities
- To keep the chamber currents stable

GIF in a nutshell



GIF gives us:

- 0.66 MeV gamma (650 GBq 137Cs)
- x30 acceleration factor in comparison to LHC
- Similar irradiation to the one expected for the RPCs in LHC

We have there:

- 6 CMS RPC double gaps (DG)
- 1 volume/hour (nominal gas mixture)
- High voltage supply
- Gaps' operation conditions and performance monitoring

Gas System



Parameters monitoring

- Control and monitoring of Environmental RH & temperature
- Source status
- Gas mixture quality
 - Composition
 - humidity before & after the purifiers
- Chambers properties and performance
 - Currents
 - Resistivity

GC/MS & HF station





Gas Chromatography

Allows quantifying ppm of impurities
3 columns – allowing separation for different type of impurities

Mass Spectrometry

• Identification of the impurities

F- ions Concentration

Based on bubbling the gas mixture and measuring of the pH of the solution

Purifiers verified at GIF





Present purifiers configuration





Filtering Capacity of Molecular Sieves

<u>Mol Sieves</u>: filter as they should H_2O (capacity ~ 150 g(H2O)/kg(MolSieve)) + filter some extra impurities + Absorb part of the RPC mix (need conditioning)



Currently 90% of Purif1 is MS3Å:

≻adsorbs water

▶ is not adsorbing $C_2H_2F_4$

≻is not effective wrt other impurities

Possible alternative is MS4Å:

- •Is not adsorbing a lot of $C_2H_2F_4$
- •Quite effective

•Release Ar (if used during regeneration)



However, many other impurities are removed (for a certain time equivalent to ~2000 volume change in the purifier cartridge)



Conditioning phase:

when a mixture component is also absorbed \rightarrow Some purifiers (see example of mol.sieve 5 Å) need a preparation time (conditioning phase) because at start-up they absorb a mixture component (in the example the $C_2H_2F_4$ is absorbed and as a result the i C_4H_{10} concentration increase).



Argon desorption from Mol. Sieve 4Å

Mol. Sieve 4Å is releasing Ar for long time after regeneration:

It can be regenerated pumping down

Alternative gas for filling during Stand-by to be found (N_2 , process gas, ...to be tested).







Filtering Capacity of R11(Cu catalyst) and R12 (Cu-Zn catalyst)

Filter as they should O_2 (capacity ~ 5 g(O2)/kg(catalyst)) + H₂O (capacity ~ 50 g(H₂O)/kg(catalyst)). R11 filters additional impurities, R12 does not and it enhance an extra component



R12: basically no extra component filtered comp. #3 is even enhanced



Filtering Capacity of Ni-Al₂O₃ catalyst



Component #2 (not present in return mixture) is strongly enhanced After a short stop, the catalyst is releasing important concentration of extracomponents (in the plot they can be compared with the SF_6 signal)



Systematic understanding of a set of purifiers vs some impurities

	Conditioning (volume change)	Main component filtered	Saturation for main component filtered (g (O ₂ /H ₂ O)/kg)	(1) CH ₄	(3) C ₂ H ₂ F ₂	(5) CH ₂ F ₂	(6) C ₂ HF ₃	(7) C ₂ H ₃ F ₃	(8) C ₃ H ₆
MS3A	3	H ₂ O	140	Unch.	Unch.	Unch.	Unch.	Unch.	Unch.
MS4A	10	H ₂ O	170	Unch.	Unch.	Rem.	Unch.	Rem.	Rem.
MS5A	50	H ₂ O	130	Unch.	Back after 1000 vol change	Rem.	Rem.	Rem.	Rem.
Cu R11	20	O ₂ /H ₂ O	5/50	Unch.	Unch.	Unch.	Rem.	Rem.	~Rem.
Cu/Zn R12	20	O ₂ /H ₂ O	5/50	Unch.	Enhanced	Unch.	Rem.	Rem.	450 vol change
Ni Al2O3	15	O ₂ /H ₂ O	15/50	Unch.	Unch.	Unch.	Rem.	150 vol change	150 vol change
Ni SiO2	15	O ₂ /H ₂ O	15/50	Unch.	Unch.	Unch.	Rem.	Unch.	Unch.



Possible new purifiers configurations

A third purifier module will be soon available (originally foreseen to be used during high luminosity LHC run) \rightarrow possibility to redistribute the cleaning agents on three module. Objective:

≻Increase the water adsorption capacity (already at the limit, it will be a real problem if any flow increase will be needed)

≻Increase the adsorption capacity for extra-Freon and extra-Hydrocarbons (but, are they really dangerous? Concentration at 50 ppm level)

≻Increase the cycle duration for the metallic catalysts (it will avoid the necessity to substitute the material during LHC running time)





Possible new purifiers configurations



<u>Conf.3 (~ Under test at GIF)</u>:

- + each metallic catalyst has one Purifier
- + double volume for Cu and Ni
- no increase of H_2O adsorption capacity
- no increase of other Freon adsorption capacity

Expected run cycles: P1: 1.4 d P2: 7 d P3: ~7 d



<u>Conf.4 (not tested) → P1 and P2 in parallel:</u> +Double the H₂O adsorption capacity +Double the other Freon adsorption capacity -P3 cycle still critical (may be necessary to change the material during LHC running time)



Possible new purifiers configurations



Conf.5 (not tested) x 2.5 the quantity of MS5Å in P1:

- + Double the H_2O adsorption capacity
- + double the adsorption capacity for other Freons
- no change for Cu and Ni catalysts

Expected run cycles:

P1: 3.5 d P2: 3.5 d P3: ~7 d



Closed loop operation



After few days of operation, only impurities 5 and 6 are still removed.
Some extra-components show higher concentration in closed loop return wrt open mode return (as expected if not completely filtered)
RPC performances do not show any degradation



Conclusions

Few critical points in the present RPC purifiers configuration V_{out} short quale for P1 (Mol. Signal) \rightarrow no margin if a flow increase

≻Very short cycle for P1 (Mol. Sieves) → no margin if a flow increase will be needed

≻Is there a limit for the maximum number of regeneration for the metallic catalysts? The present used limits comes from max number of tested cycles during previous test at GIF (and recommendations from producer)

≻Very limited action on extra-Freon/Hydrocarbons, but are they really dangerous?

≻Possible improvement from new purifier configuration:



> but MS4Å desorbs Argon (if used during regeneration). Alternative candidate for fill during Stand-by phase to be found (N₂ or process gas) or no Stand-by (i.e. directly to Preparation for Run after Cool down) or