

A new gas system for the Würzburg cosmic ray facility

Pipe Studies

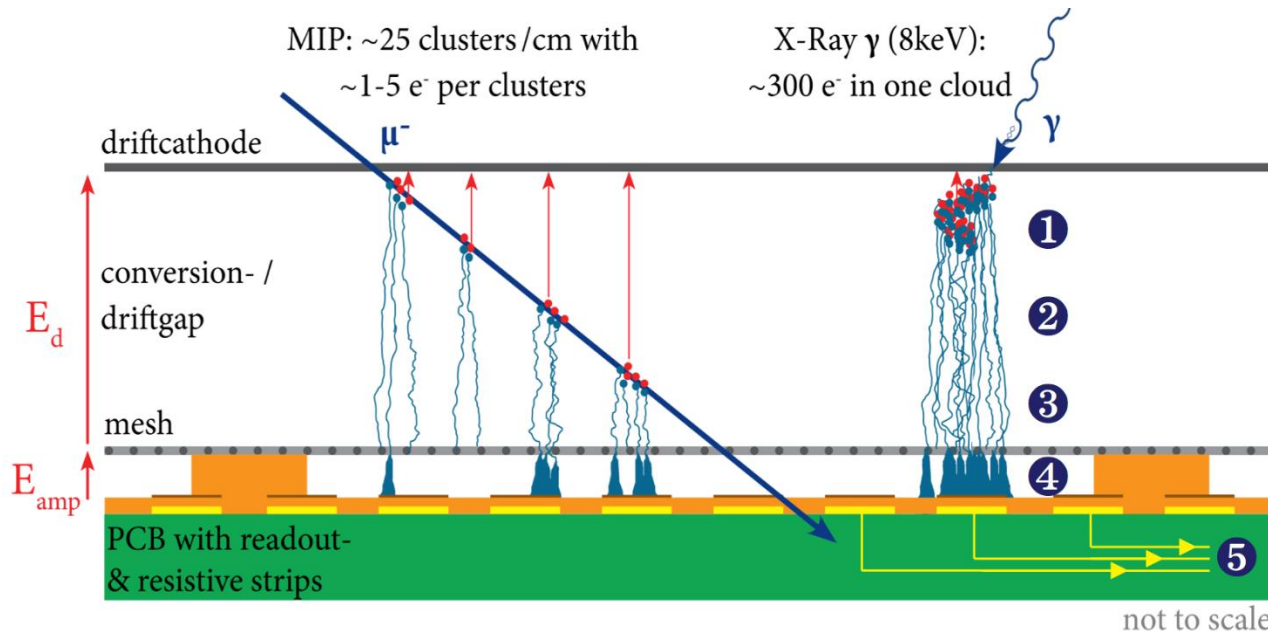
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RD51 Collaboration Meeting
2019/10/23 CERN

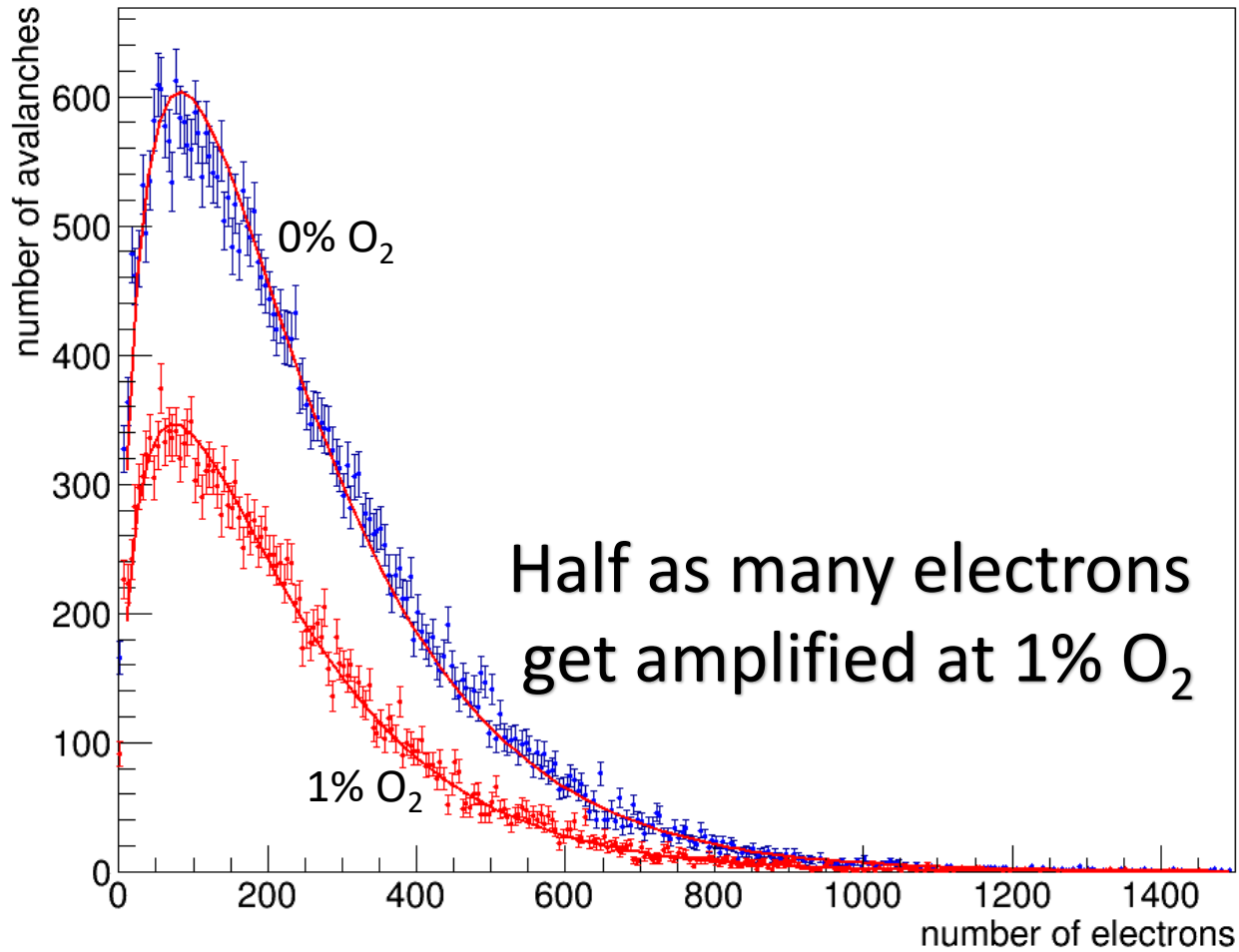
- RnD on gaseous particle detectors is being done in Würzburg
- Effects of small gas contaminations not yet fully investigated
- The new gas system will be able to regulate small concentrations of O₂ and H₂O (ppm to 1%) and measure them at the exhaust

- Both Water and Oxygen influence the detector performance
 - Oxygen is highly electronegative and free electrons can get lost due to attachment to it
 - Water influences the high-voltage stability of the detector



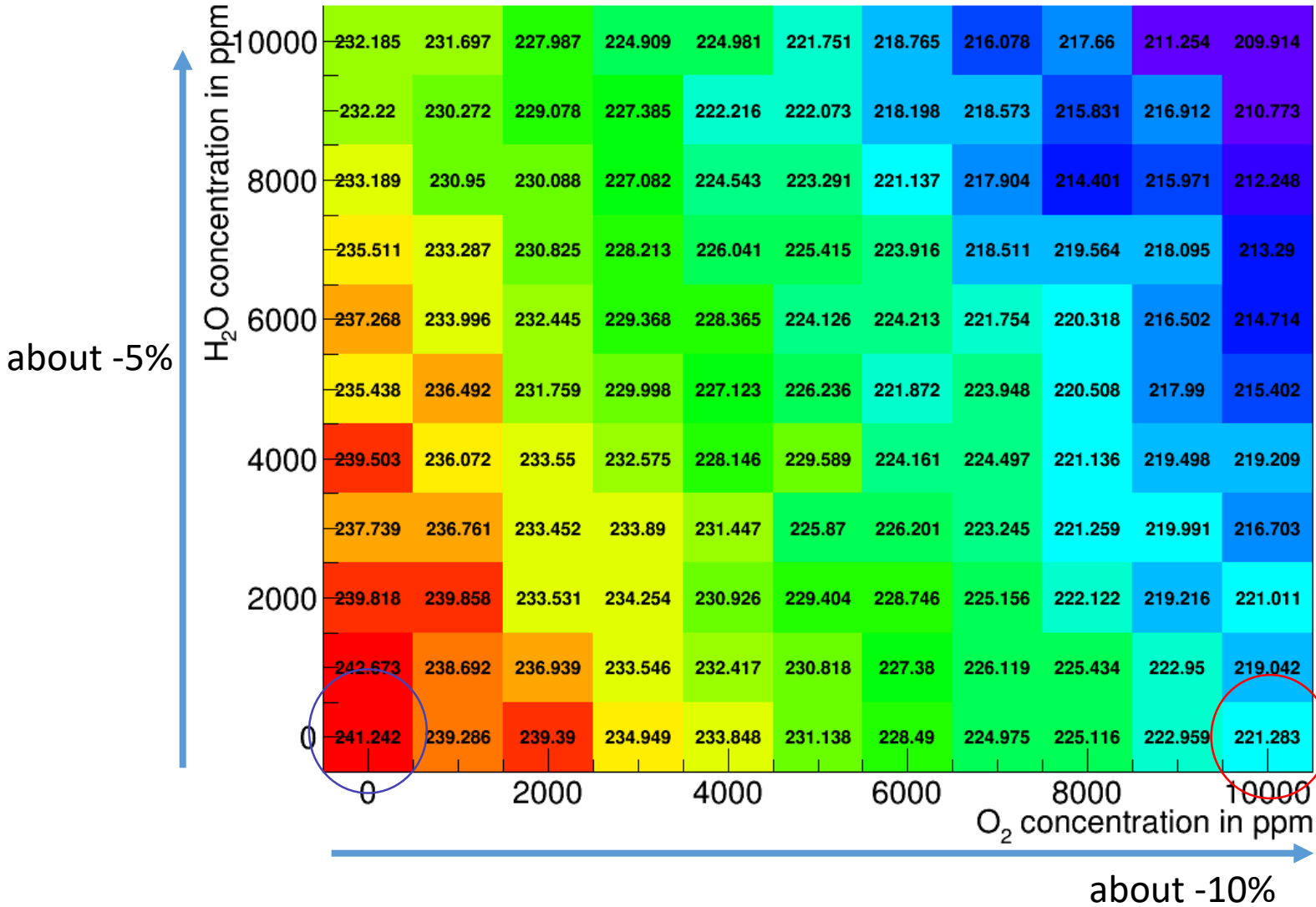
Simulations in Garfield have been done:

- Includes full Micromegas field structure
- Base gas mixture is 93:7 Ar:CO₂
- Penning transfer is disabled
- 5000 muon events have been simulated
- Incident muon creates primary electrons
- Every electron is tracked through the detector and amplification is simulated
- Both gain and number of amplified primaries are important values

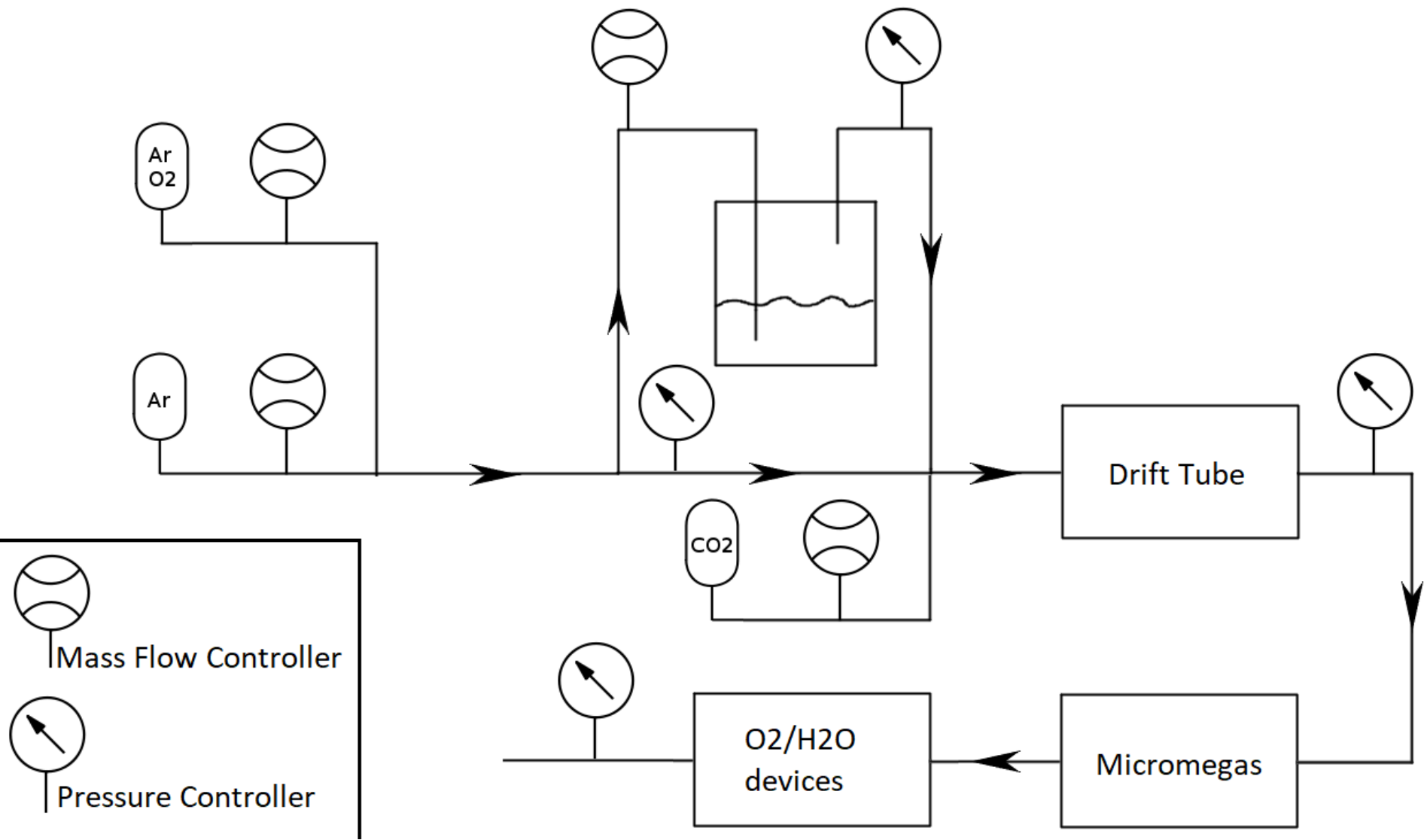


THE SIMULATION GRID (GAIN)

10000 ppm = 1%



THE GAS SYSTEM SCHEMATICS



- Systech Illinois ZR800:

- 0.1ppm – 100% O₂
- 0.2% relative uncertainty

- Systech Illinois MM400:

- 20ppm-2.4% H₂O
- 2 ppm or 0.5% uncertainty



- First tests have been done to verify the purity of the gas before any contaminations are added on purpose
- About 20 ppm O₂ and 400 ppm H₂O are already present.
- Conjecture: any remaining plastic tubing could allow exchange
- Test: remove all plastic – only metal pipes
- Result: <1 ppm O₂ and <1 ppm H₂O

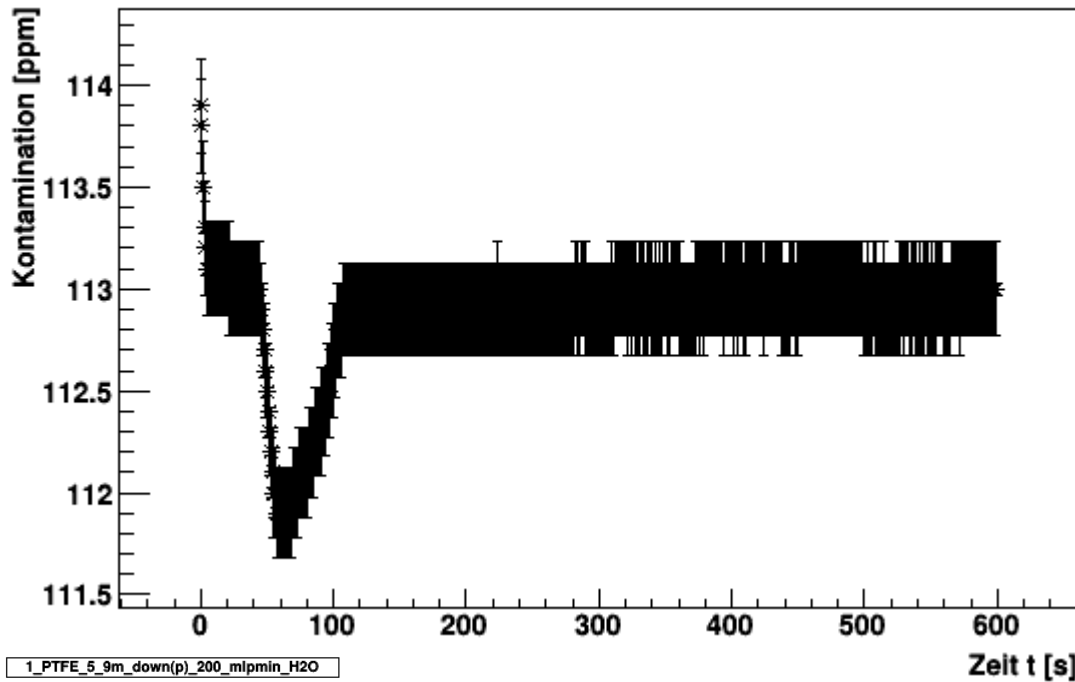
- Plastic tubing is very common in experiments using gaseous detectors
- Therefore, the question of which material to use to minimize contamination is of utmost importance
- Being able to quantify expected contamination is also important
- The following plastic tubing has been tested

Material	Inner Diameters [mm]	Wall thickness [mm]
PTFE	3.7, 4.0, 4.5, 5.0	1.15, 1.0, 0.75, 0.5
PU	4	1
PVC*	4	1
PFA	4	1

*ordered, but too soft to be connected to the system

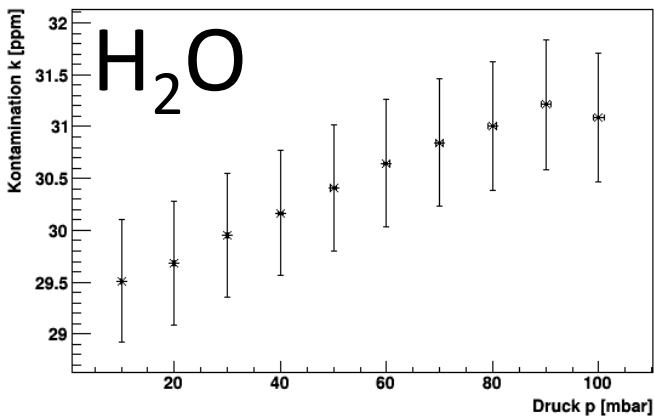
- 10 m of each type of pipe is available, cut to 1, 2, 3, 4 m (not everything has been measured)
- For each piece, uncontaminated gas is flushed at different flows and pressures:
 - Flushing at 200 ml_s/min for a few hours
 - 200...20 ml_s/min in steps of 20 ml_s/min at 50mbar_g
 - 20...200 ml_s/min in steps of 20 ml_s/min at 50mbar_g
 - 10...100 mbar_g in steps of 10 mbar_g at 200 ml_s/min
- Readings of both Oxygen and Water concentrations are taken every second
- Following measurements have been taken by a BSc. student

2_PTFE_5_9m_down(p)200mlpmin_80mbar_O2

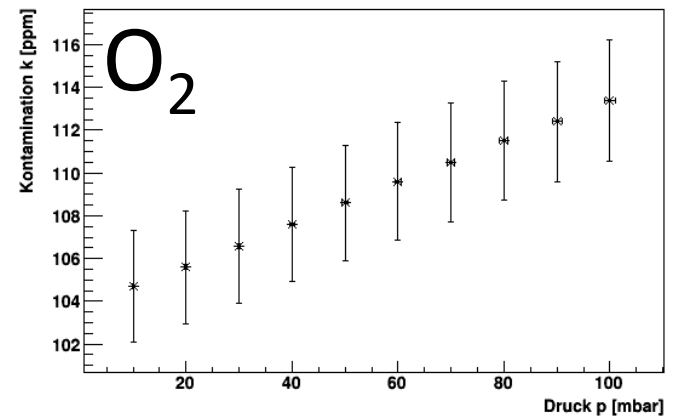


- Pressure change in the measurement devices can be seen
- Guess: Change in calibration factor leads to slight change of output

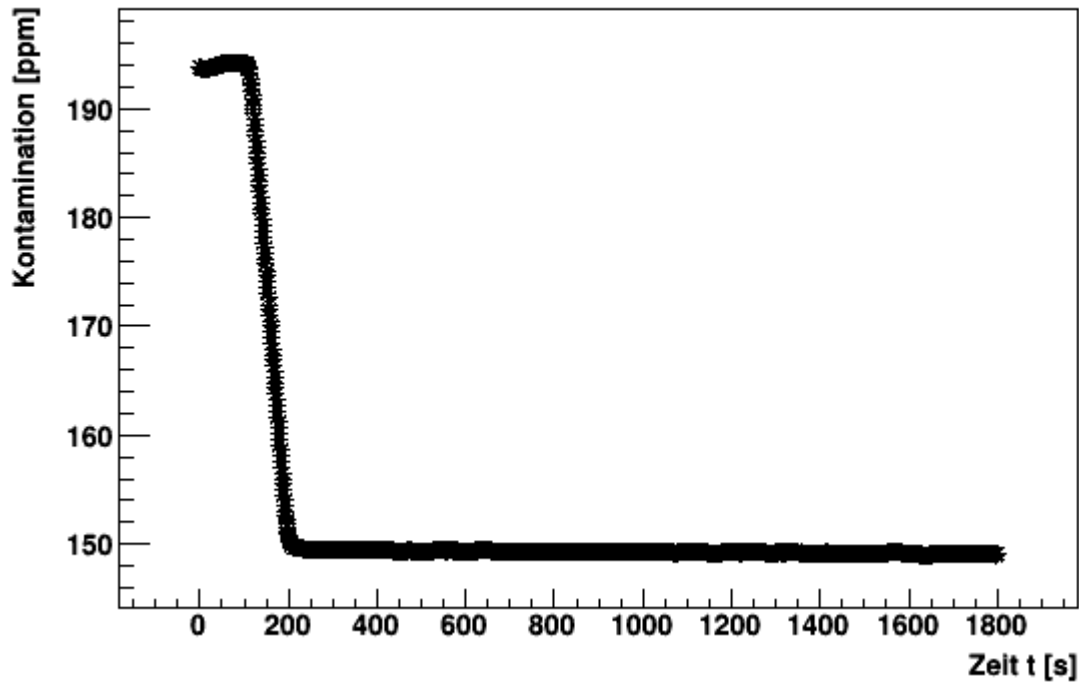
1_PTFE_5_9m_down(p)_200_mlpmin_H2O



1_PTFE_5_9m_down(p)_200_mlpmin_O2



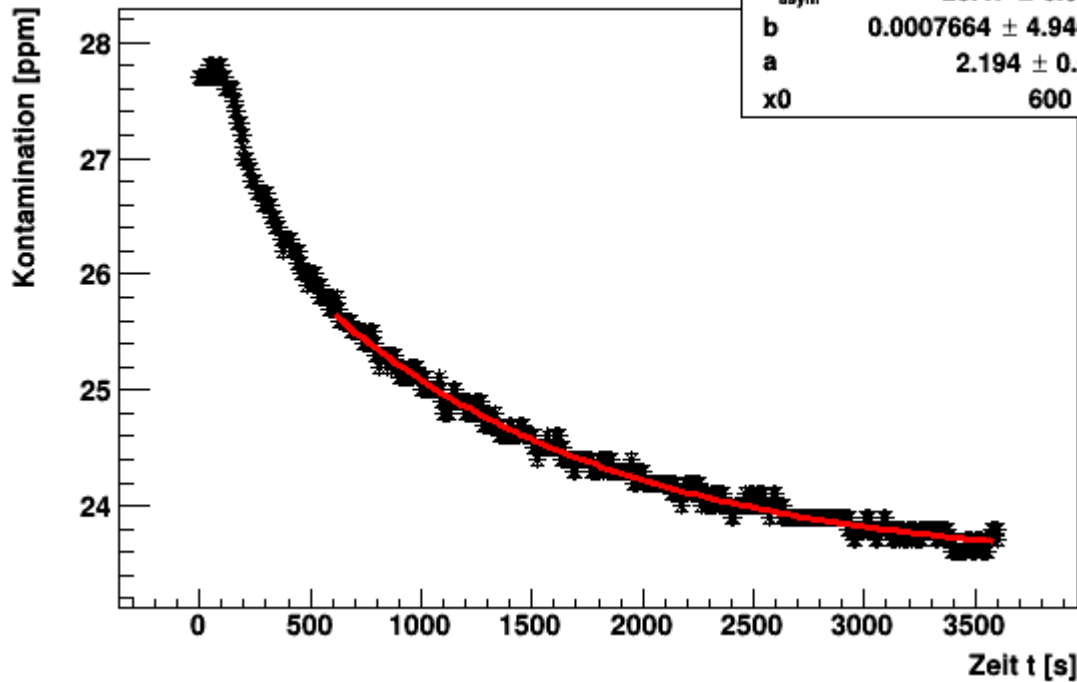
2_PTFE_4_10m_up(f)80mlpmin_50mbar_O2



- Concentration change almost instant
- After that, the concentration stays constant
- Just reading out the last measurement is enough

1_PTFE_37_9m_up(f)100mlpmin_50mbar_H2O

χ^2 / ndf	3864 / 2997
y_{asym}	23.47 ± 0.005396
b	$0.0007664 \pm 4.946e-06$
a	2.194 ± 0.00447
x_0	600 ± 0

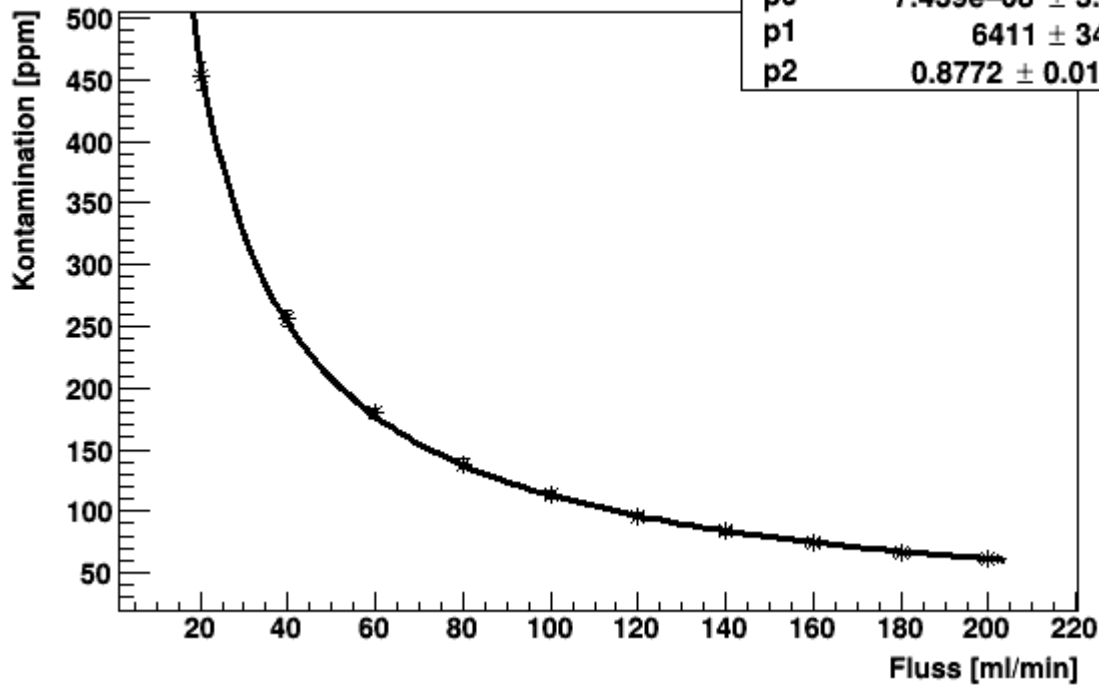


$$c = y_{\text{asym}} - a \cdot e^{b \cdot (t - x_0)}$$

- Changes are taking a long time
- Fit appropriate, with x_0 fixed
- It seems like there are both a short and a long change rate
- Therefore, the first part is taken out of the fit

1_PTFE_4_9m_up(f)_50_mbar_O2

χ^2 / ndf	1.714 / 7
p0	7.439e-08 ± 3.495
p1	6411 ± 348.6
p2	0.8772 ± 0.01192

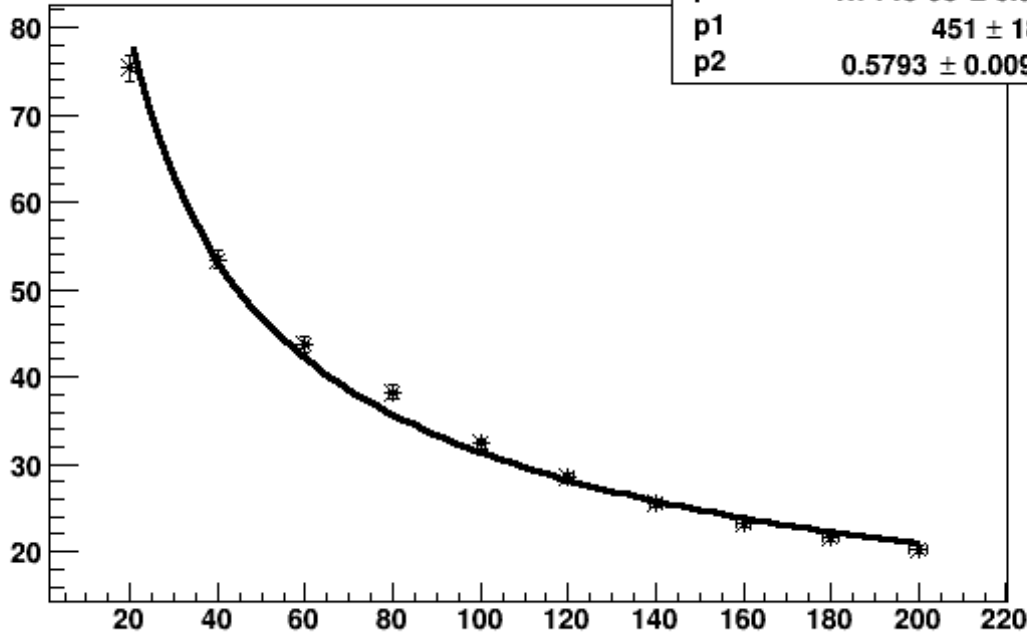


$$c = p_0 + p_2 \cdot x^{-p_1}$$

- Guess: contamination depends on time in the pipe (about linear)
- Fit a power function
- Check for constant offset (other sources!)
- Scale factor p_2 is a property of the pipe:
 - Material
 - Length
 - Wall thickness

1_PTFE_45_9m_up(f)_50_mbar_H2O

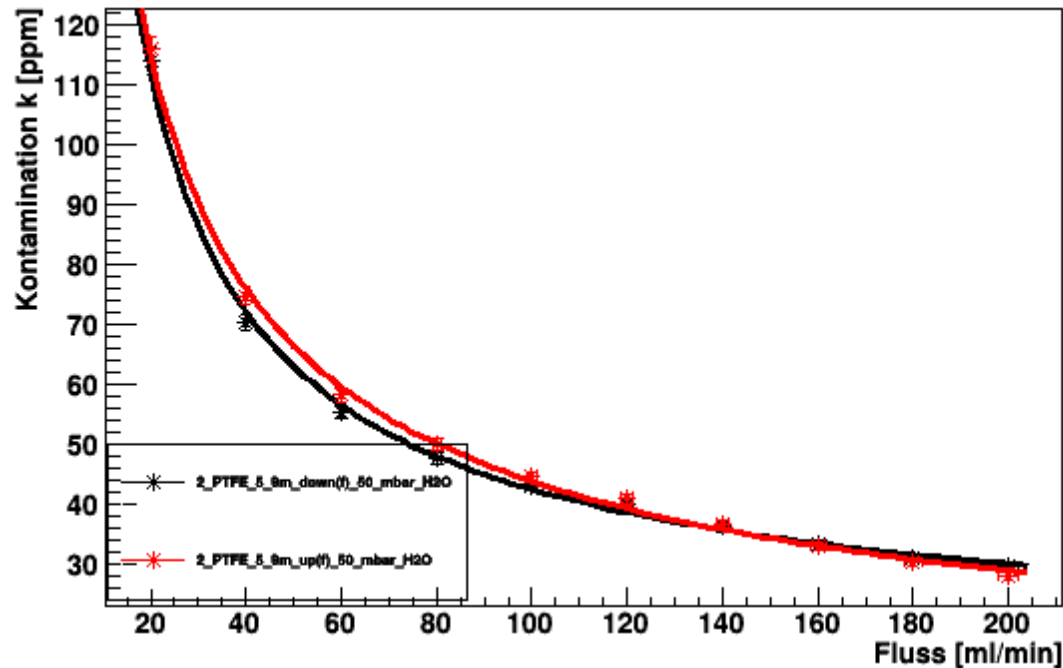
χ^2 / ndf	29.73 / 7
p0	1.714e-08 ± 0.3104
p1	451 ± 18.75
p2	0.5793 ± 0.009108



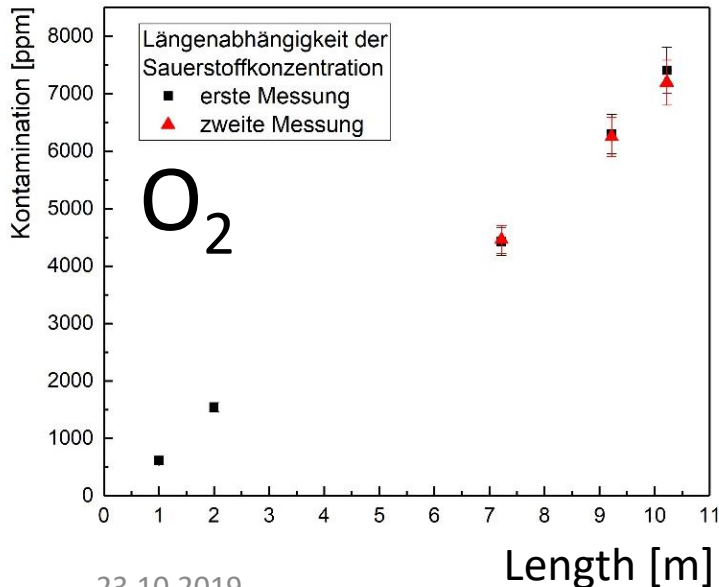
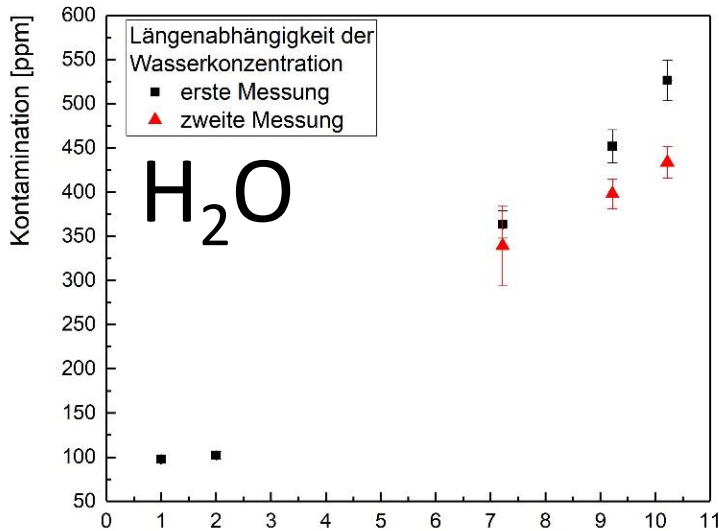
$$c = p_0 + p_2 \cdot x^{-p_1}$$

- Fit worse than in the case of O₂
- Square root dependence, not linear!
- Hint at different mechanism for contamination
- Comparison between two measurements of the same pipe might be interesting

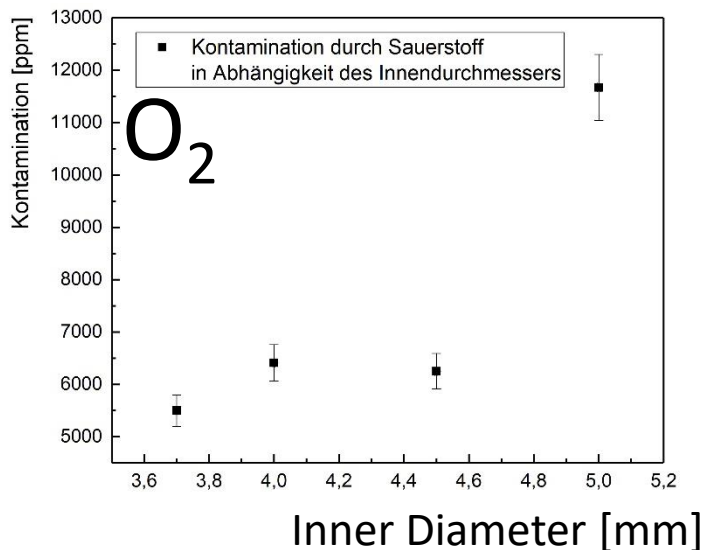
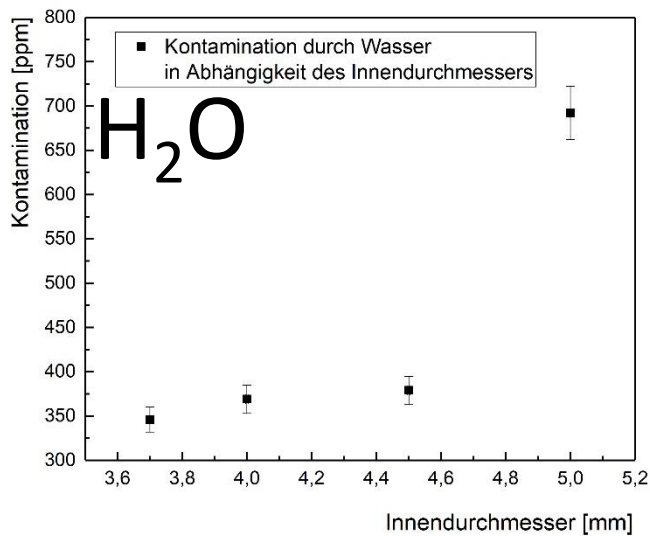
H₂O comparison



- Same pipe in both curves
- Measurements one right after the another
- Black is going down, red is going up
- Because of less influence from open air, the up(flow) measurement will be used further
- Problem: Air humidity in the room is neither controlled nor tracked!



- Black and red are:
 - Same pipe (PTFE, 6x4mm)
 - p_2 extracted from up(flow) measurements
 - Two different days (each flushing, down(flow), up(flow), down(pressure))
- O_2 is almost perfectly repeatable
- H_2O is not, air humidity logging is being worked on



- Inner diameter is not too important for contamination if small enough
- Seems like a plateau, with the contamination decreasing towards the lower diameters and increasing towards the higher
- Those are all 6mm outer diameter tubing variations available, 2 metric, and 2 imperial

Material	p_2 of O ₂	p_2 of H ₂ O
PTFE	6400±350	340±15
PFU	450±255	800±240
PFA*	3660±190	770±110

*PFA measured at 6m and scaled by a factor 1.5

- PTFE and PFA show similar behaviour
 - Better at H₂O than at O₂ by an order of magnitude
 - PFA has a less pronounced difference between the two contaminations
- PFU is susceptible to H₂O but less so to O₂
- Choice of pipe depends on the contamination you want to reduce, but the best choice would be metal!

- The new gas system in the Würzburg cosmic ray facility is now ready for precise measurement and is producing first results
- This prompted an investigation into the permeability of piping of different material and thickness
- The first results have been shown here
- The studies have a general relevance for all gaseous detector systems
- Goal: measuring the effect of O_2 and H_2O on detector performance

A big thank you to our
scientific workshop and
thank you for your
attention!

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