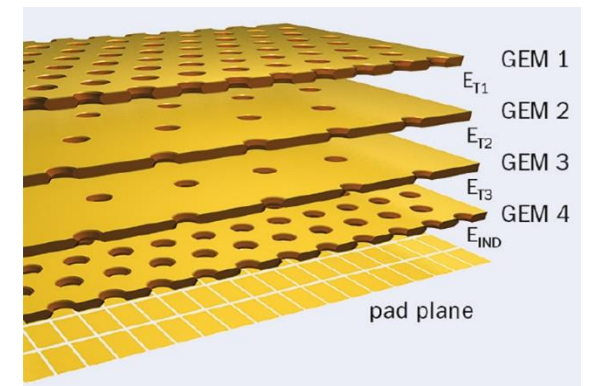


Some features of the GEM -TPC prototype operation in Ne+CF4

C. Garabatos, V. Peskov

Behalf of the ALICE TPC upgrade team



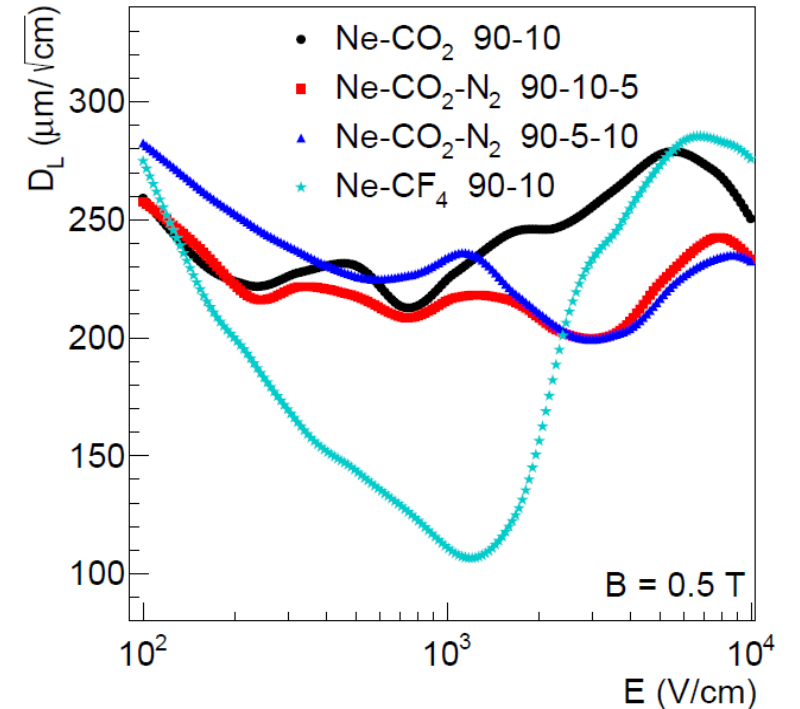
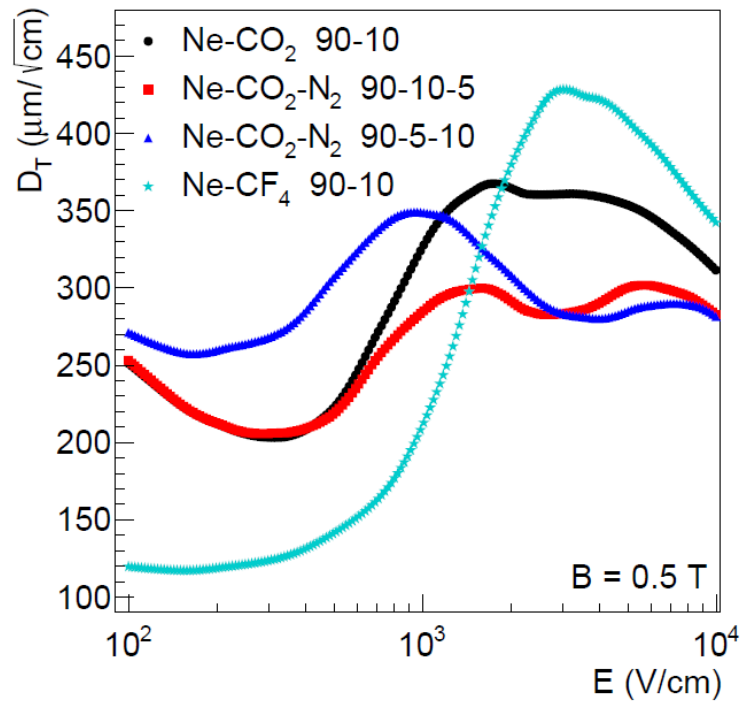
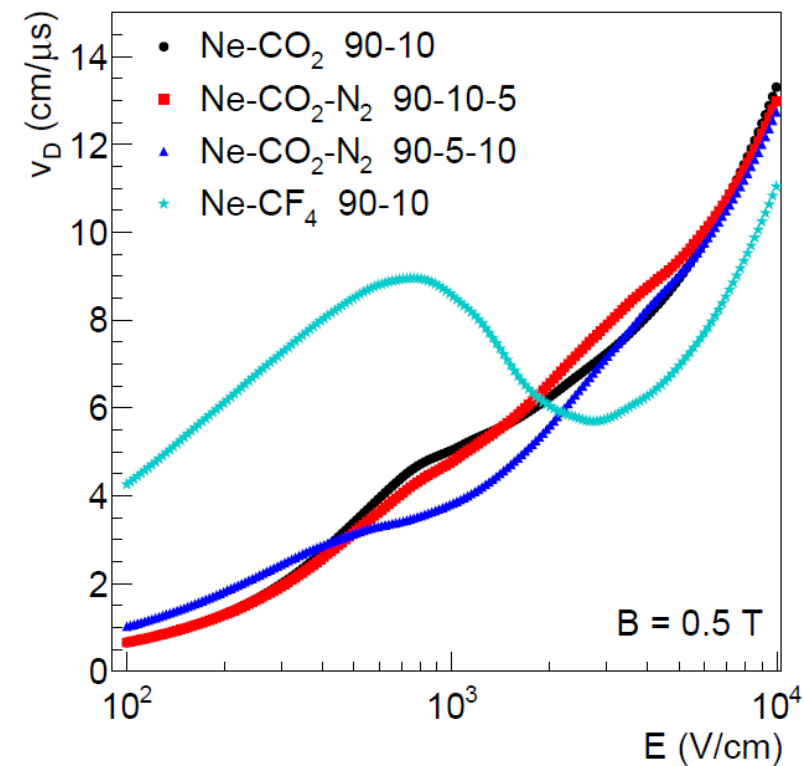
This message, we received among others from one of our s-PHENIX colleagues, stimulated our presentation:

“...I attach a plot we generated during the ALICE R&D. We tested 90% Ne + 10% CF₄. We found that there is something like resonant capture of electrons on CF₄. The plot was generated for a 2-GEM MMG chamber, varying the field between the two GEMs. Our colleagues in Europe verified that the same happens with a 4-GEM chamber. If you run a 4-GEM setup with the high transfer fields that are needed for best IBF suppression (4kV/cm) the chamber gain is lower by at least a factor of ~10 from that at a transfer field of ~1 - 1.5 kV/cm. Anyhow I'd like to raise this as a point for discussion of which gas really is the best for the sphenix tpc..”



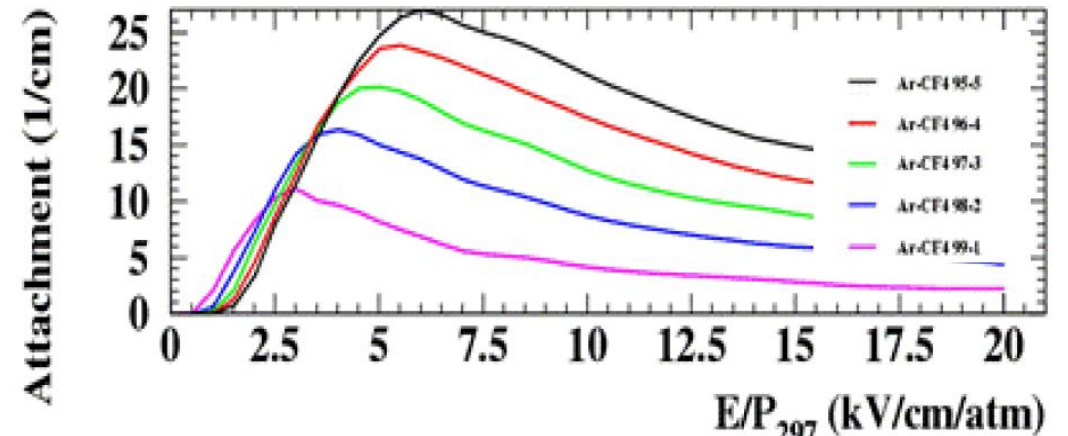
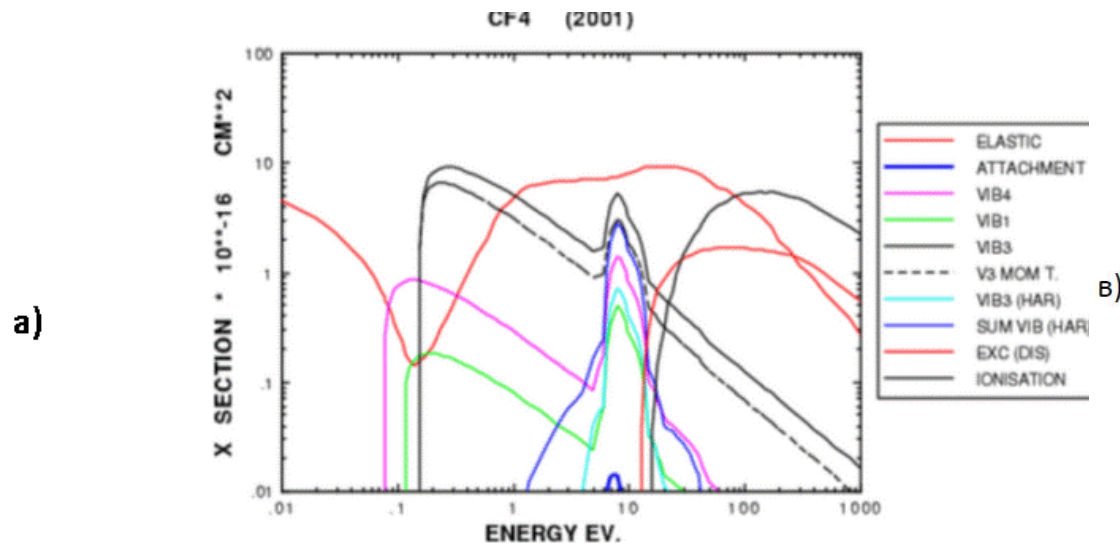
We, thus, think that it will be useful to give more details from these ALICE studies

Reminder: why some people think that CF₄ could be an interesting alternative to Ne+CO₂+N₂?



...because in this gas in some voltage interval electrons have faster drift velocity and lower diffusion

The challenge, however is the CF4 electronegativity



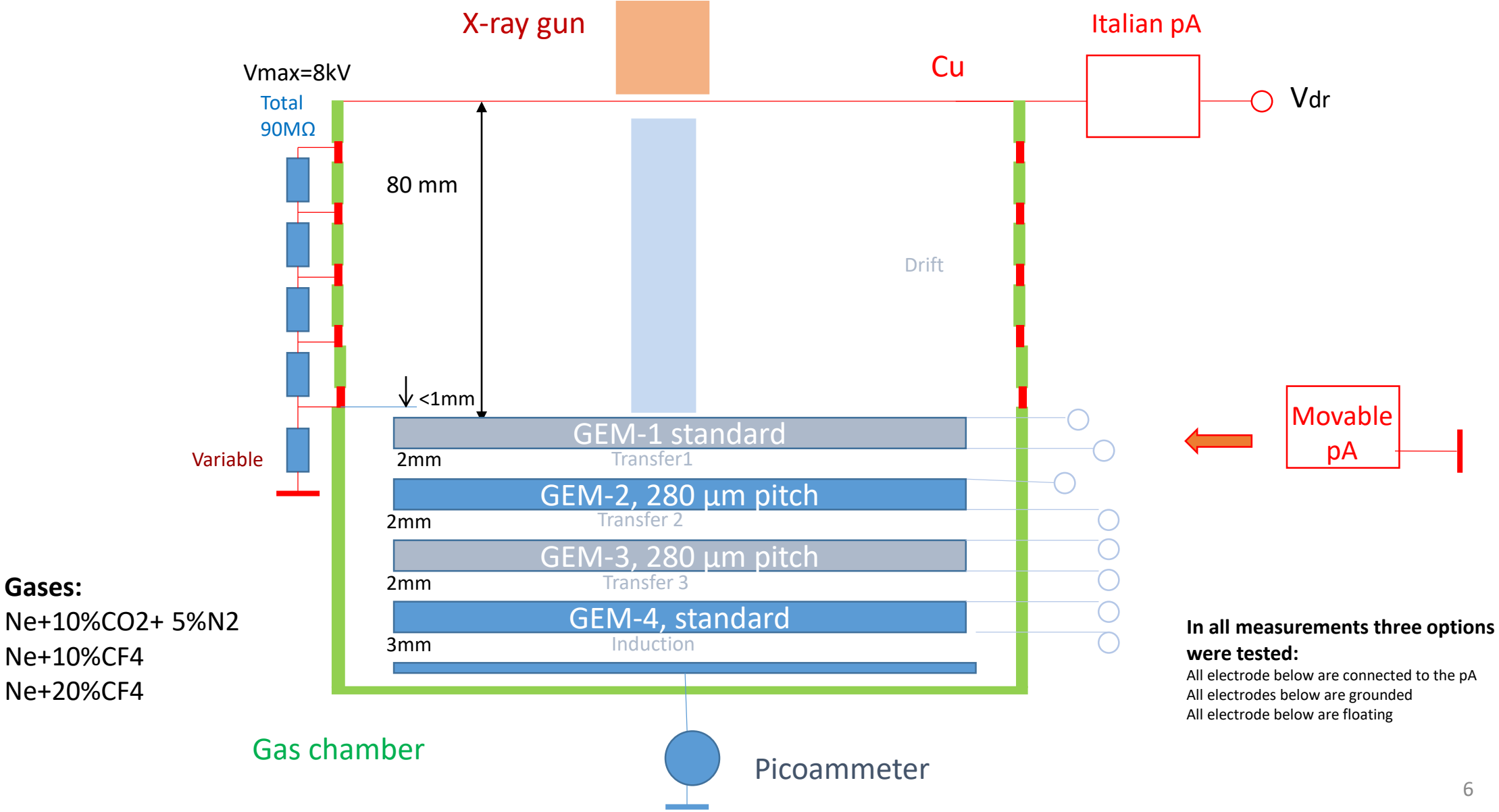
ALICE TPC upgrade group made some pilot studies of a quadruple GEM operation in CF4 mixtures focused mainly on the IBF issue



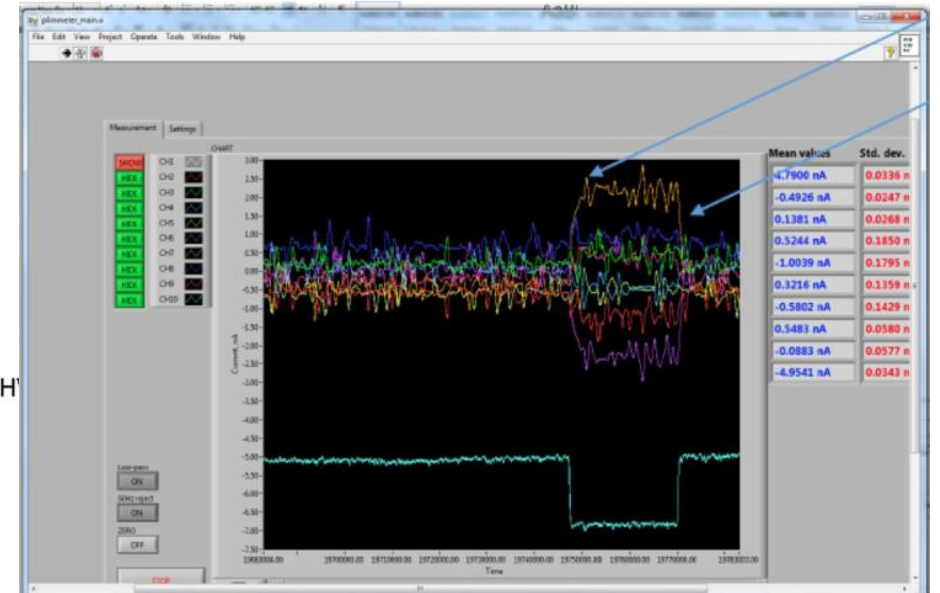
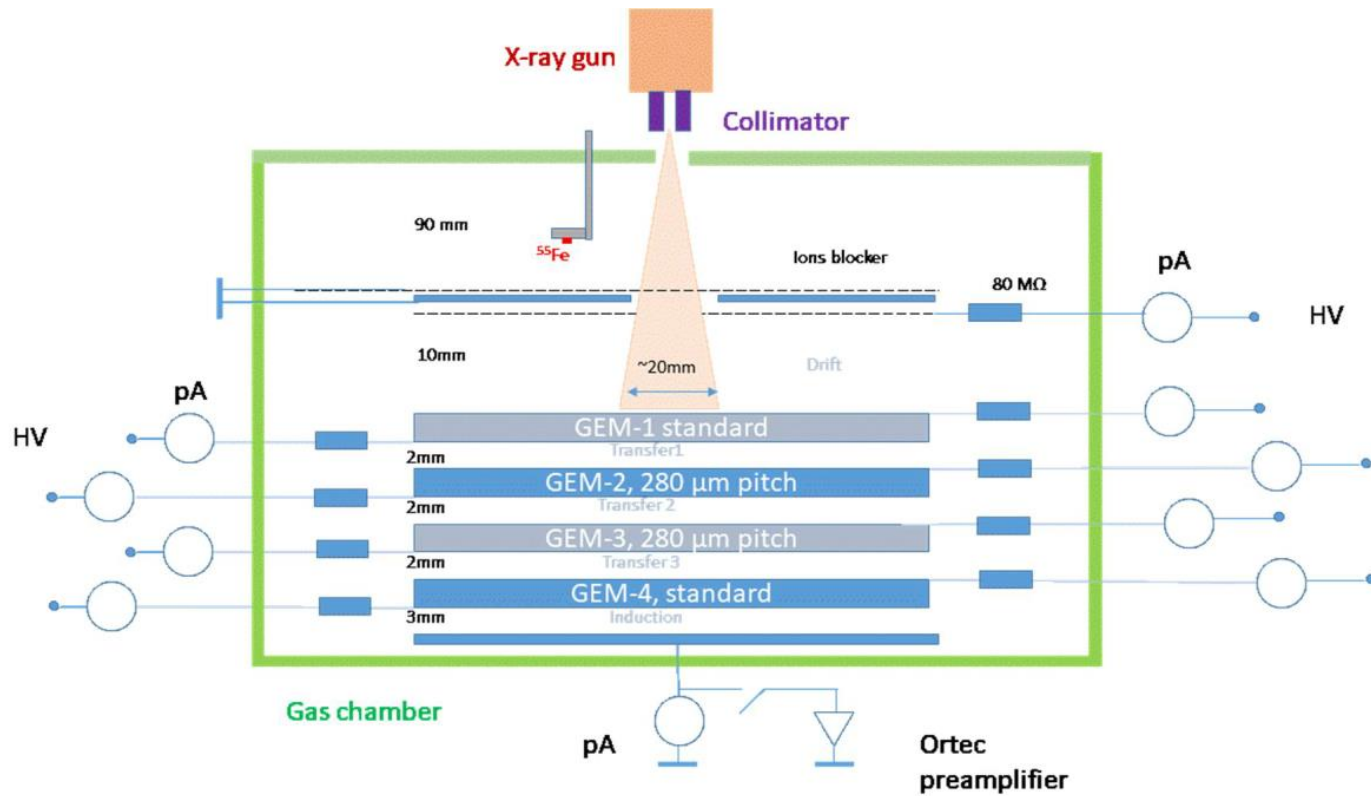
We will present some of them: comparison of quadruple GEM operation in Ne+CO₂+N₂ and Ne +CF₄ (10 and 20%)

(note: these ALICE results were never presented outside ALICE community)

Experimental setup

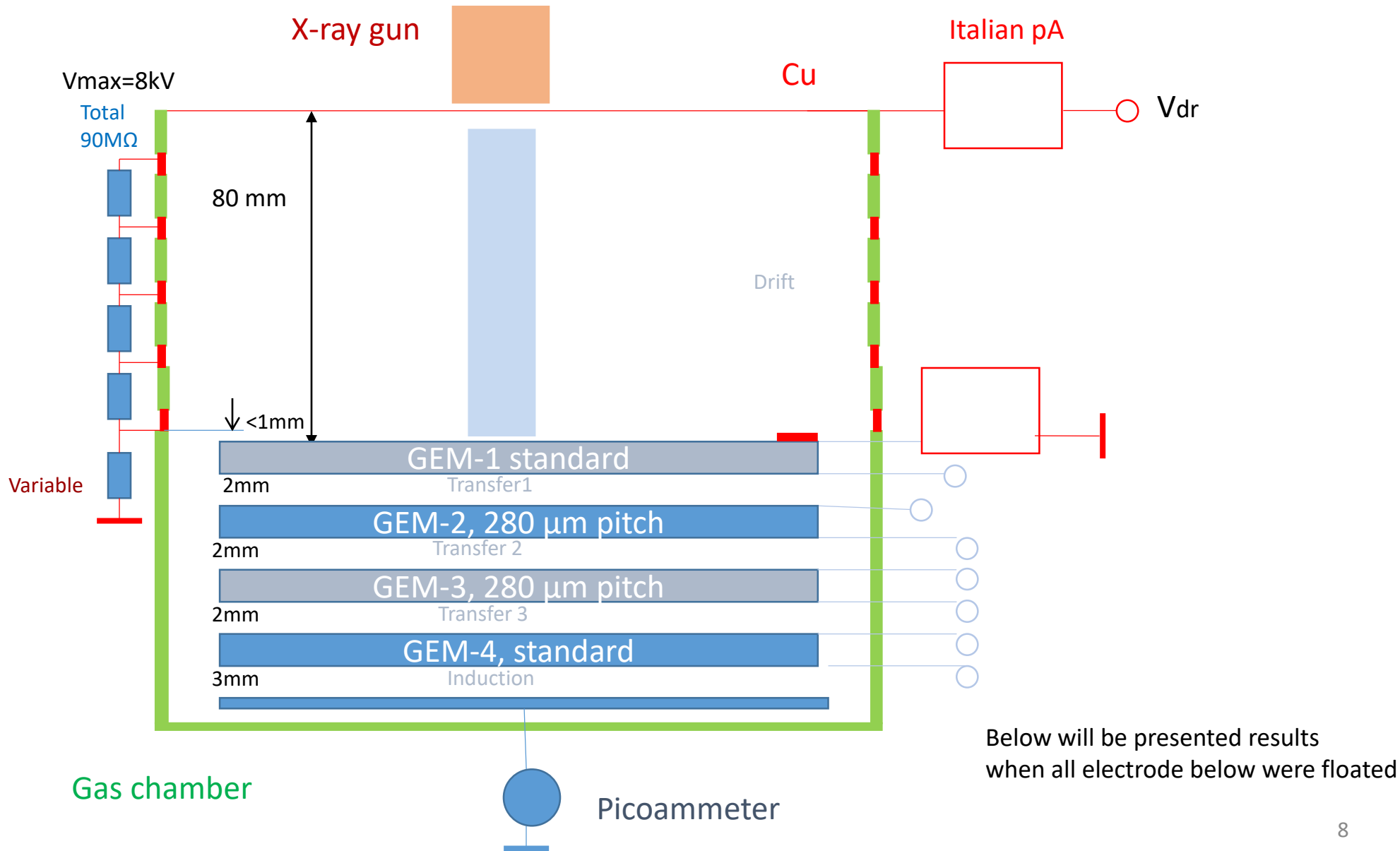


Some measurements were repeated at WIS, using a more sophisticated set-up, when one could control all GEM currents simultaneously



... but today we will show only CERN results

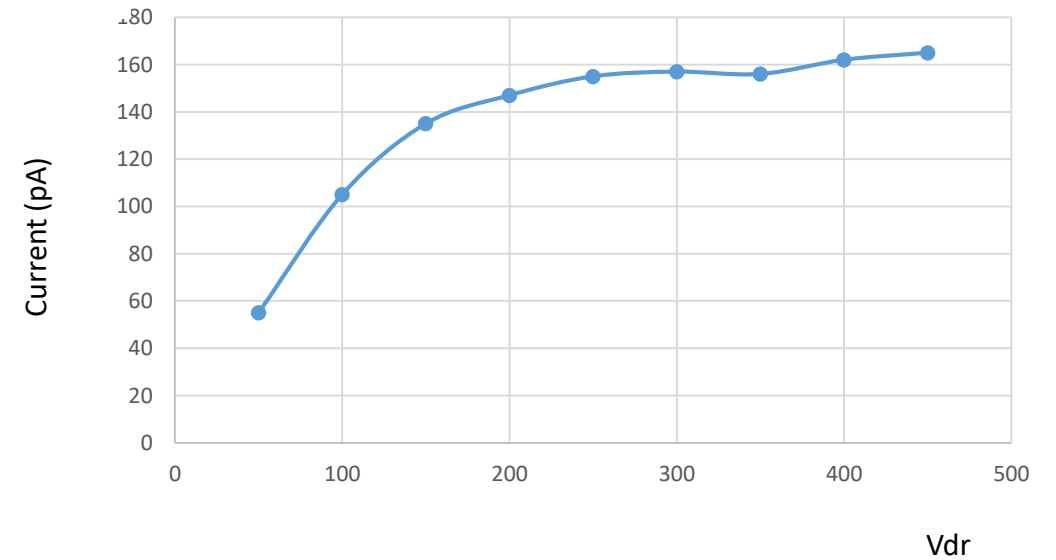
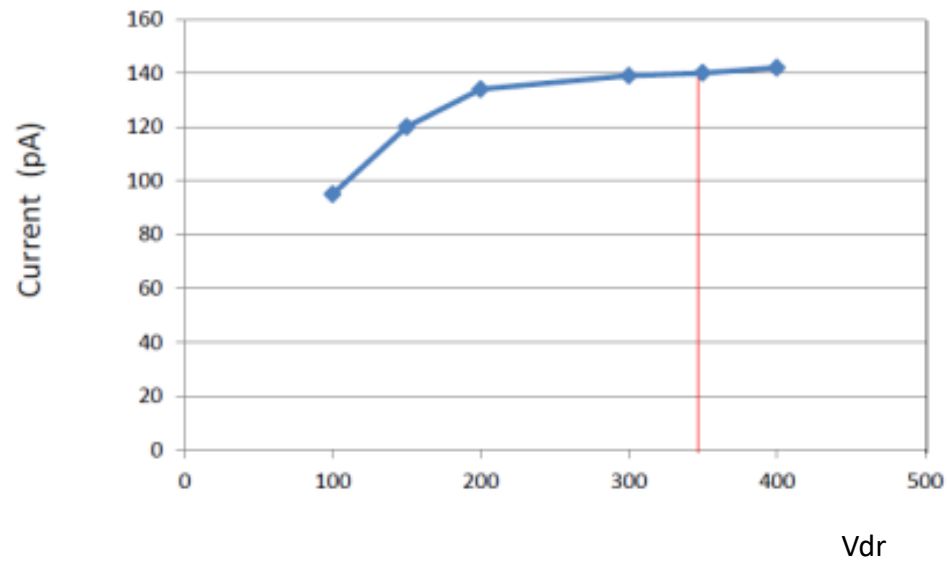
Measurements of the primary ionization current



Ne+10%CO2+5%N2

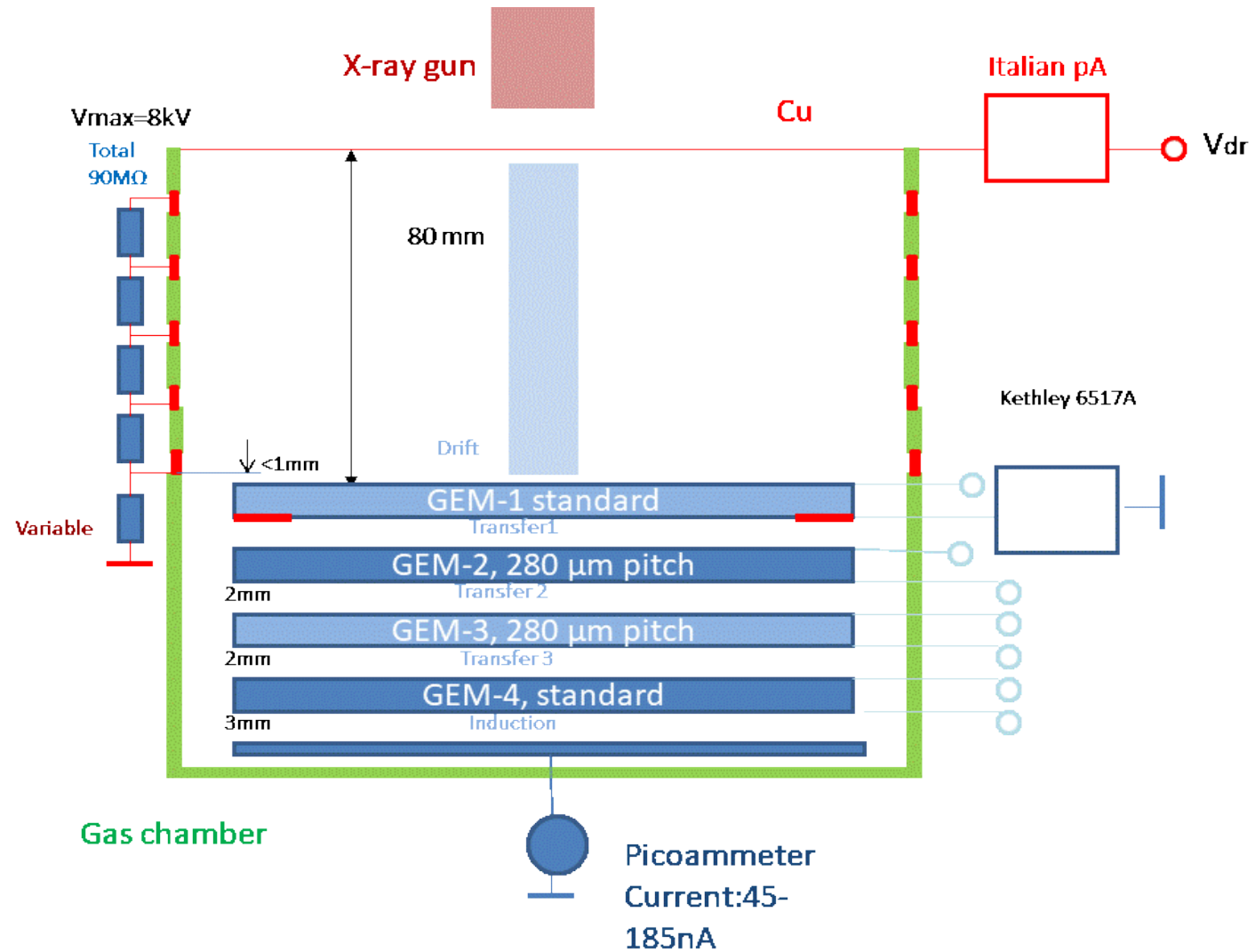
Ne+20%CF4

X-rays:
40kV, 100mkA

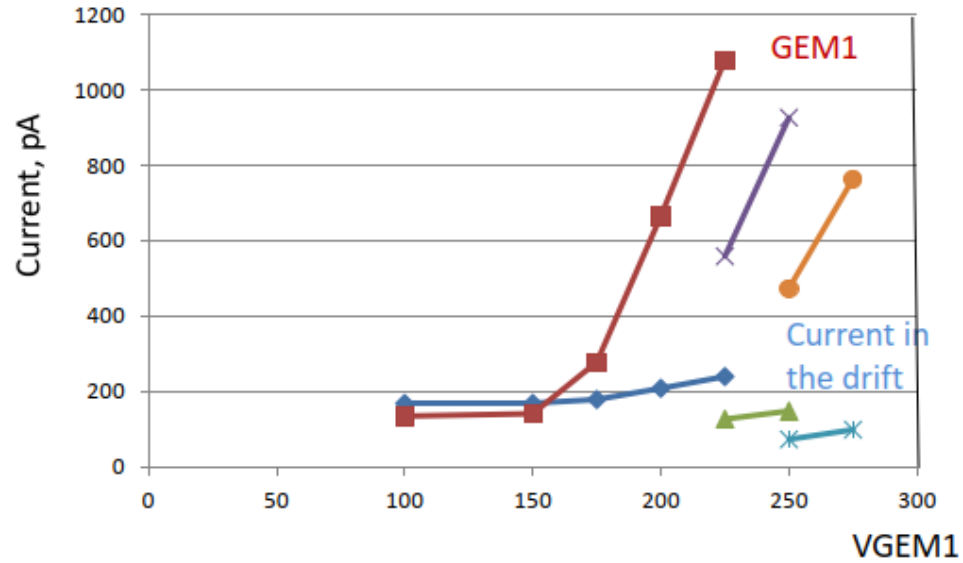


At a drift voltage of 200V/cm electron capture by CF4 is small, so it is not astonishing that the results in both gases were similar

Measurements of multiplication in first GEM



Raw data, obtained at different X-ray currents

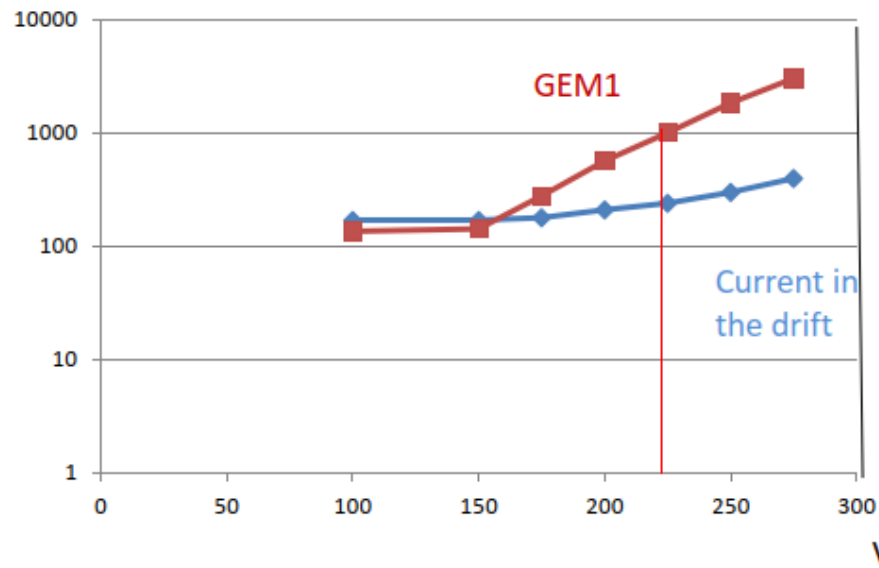


Ne+10%CO2+5%N2

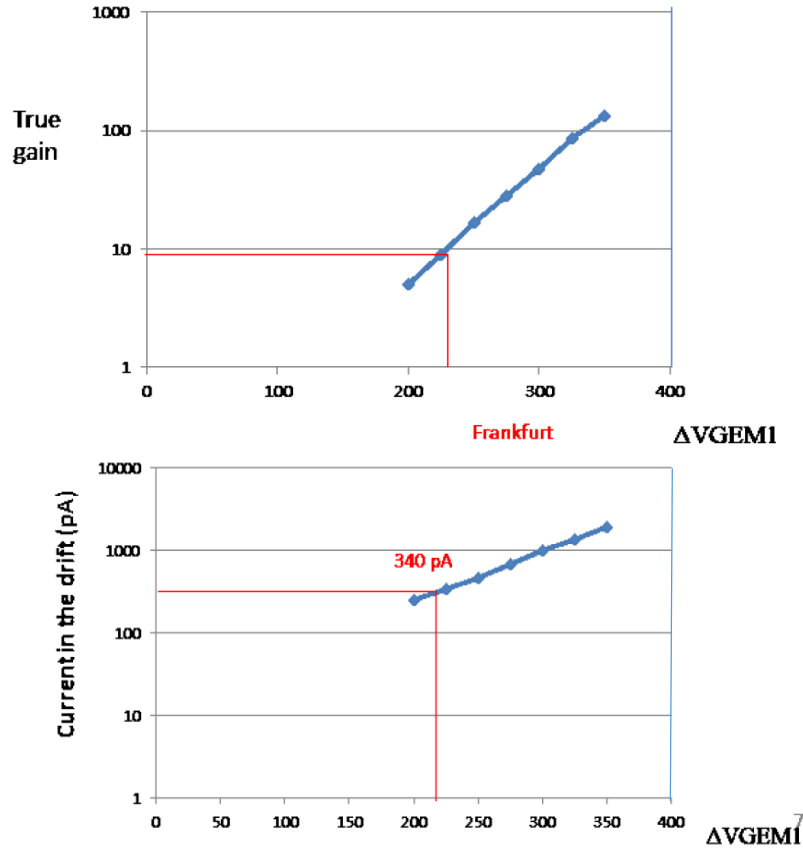
We used the max setting to measure the primary current in the drift and then, when operating with a gain the x-ray current was reduced to avoid the space charge effect (we always kept the current on the readout plate below 10nA, which is expected current at the ALICE experiment). However, presenting the data, obtaining with the gain, we often, for convenience, recalculate then to the setting **40/100** in order to see immediately the IBF.

Combined plot for **40/100**

At VGEM1=225
Gain = 7.2,
IBF≈24%



Ne+10%CO2+5%N2



$$A_1 = I_{GEM1} / I_{inj}$$

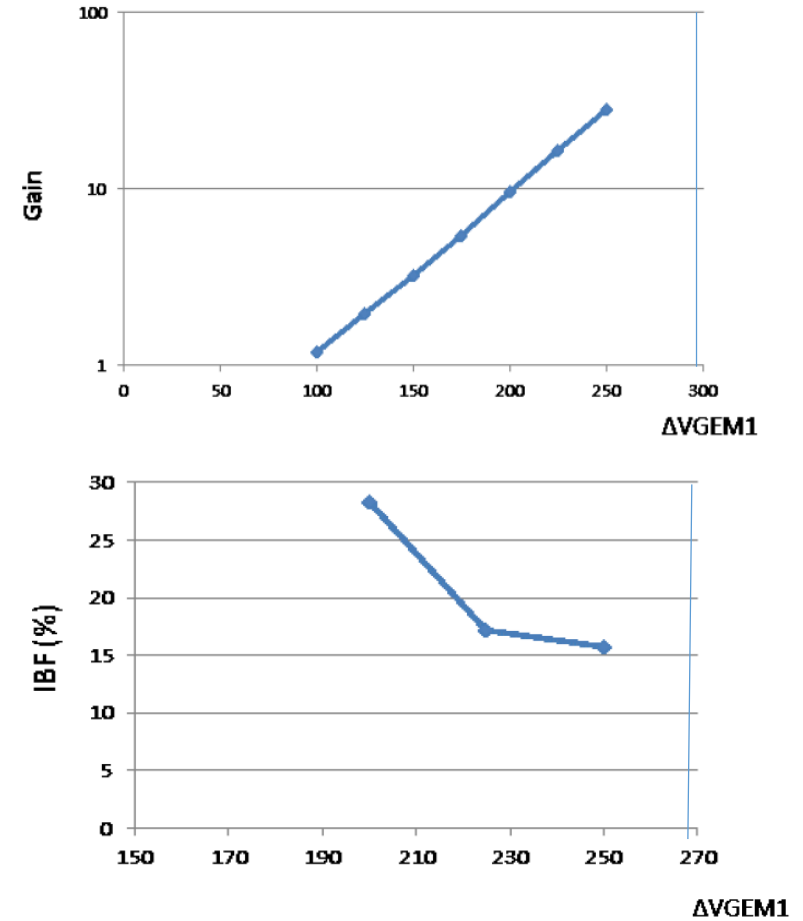
$$ABF = (I_{dr} - I_{ioniz}) / I_{GEM1}$$

X-ray setting
40/100

In a uniform field $A = \exp(\alpha)x$

Ne+20%CF4

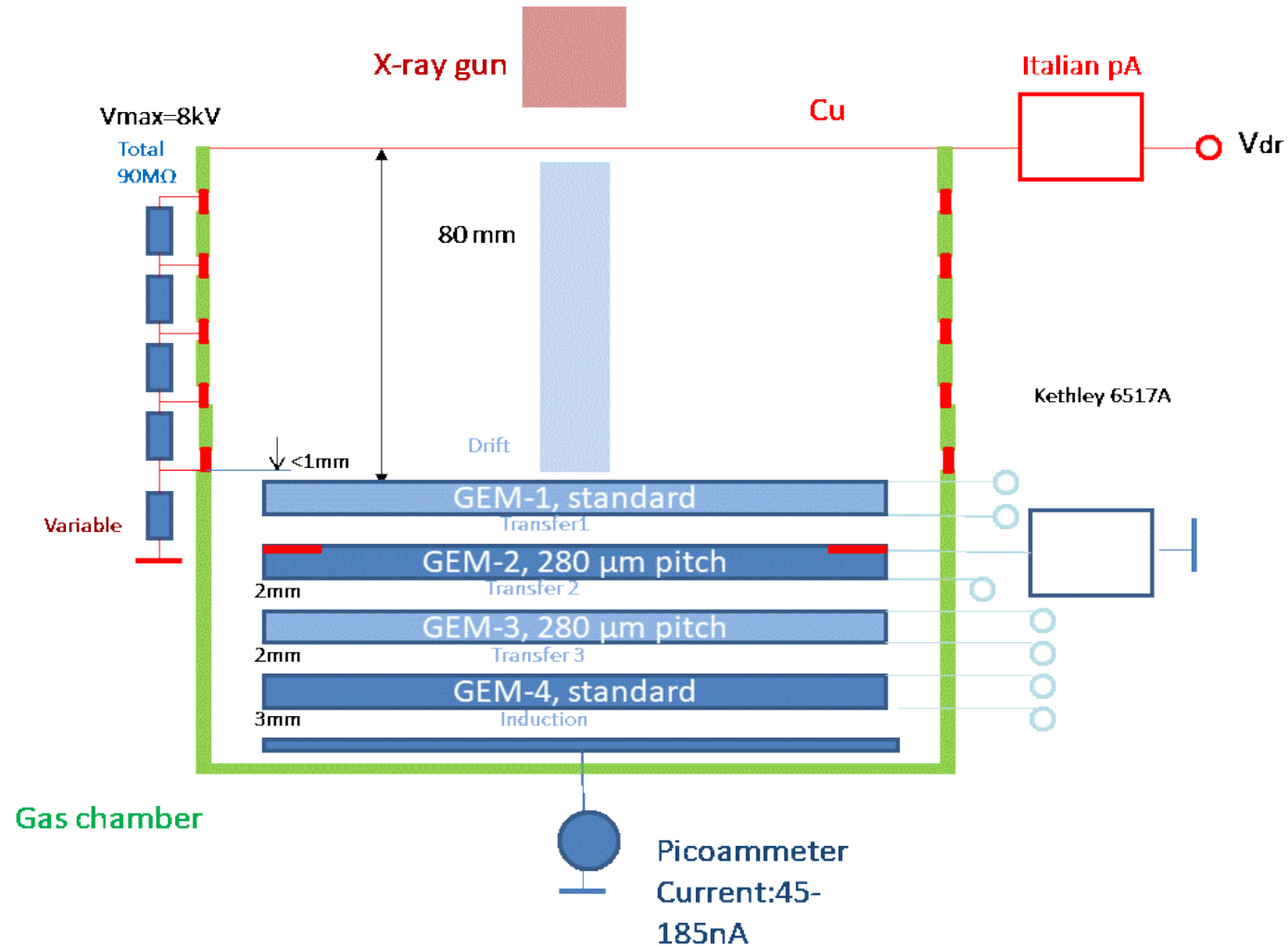
GEMs gain



$A = \exp(\alpha - \eta)x$

In both cases the dependence should be exponential and this was observed experimentally

Measurements of extraction from first GEM

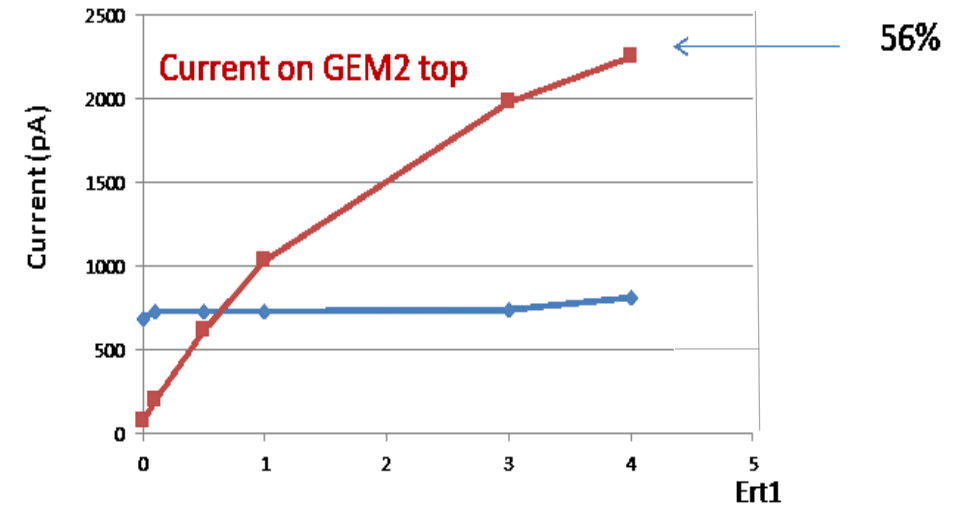
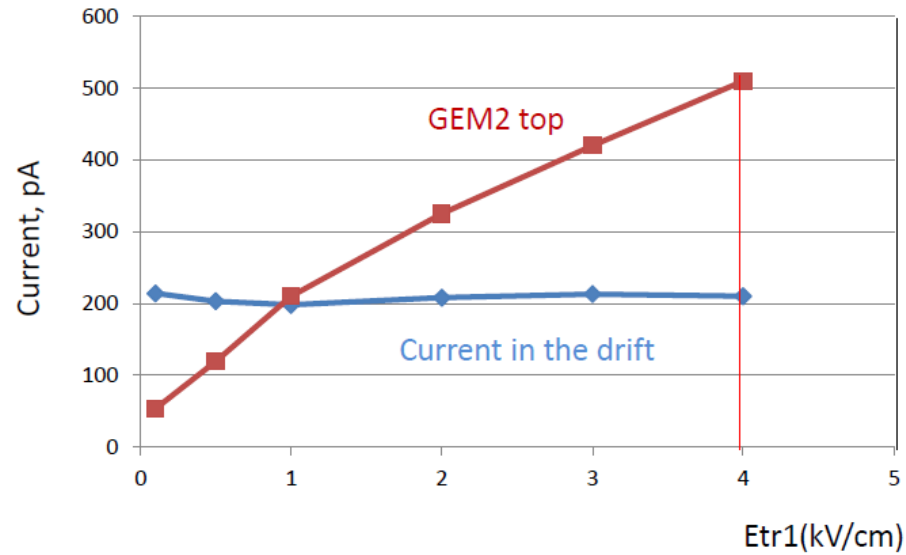


Ne+CO₂+N₂

Ne+CF₄

VGEM1=225
40/100

40/100

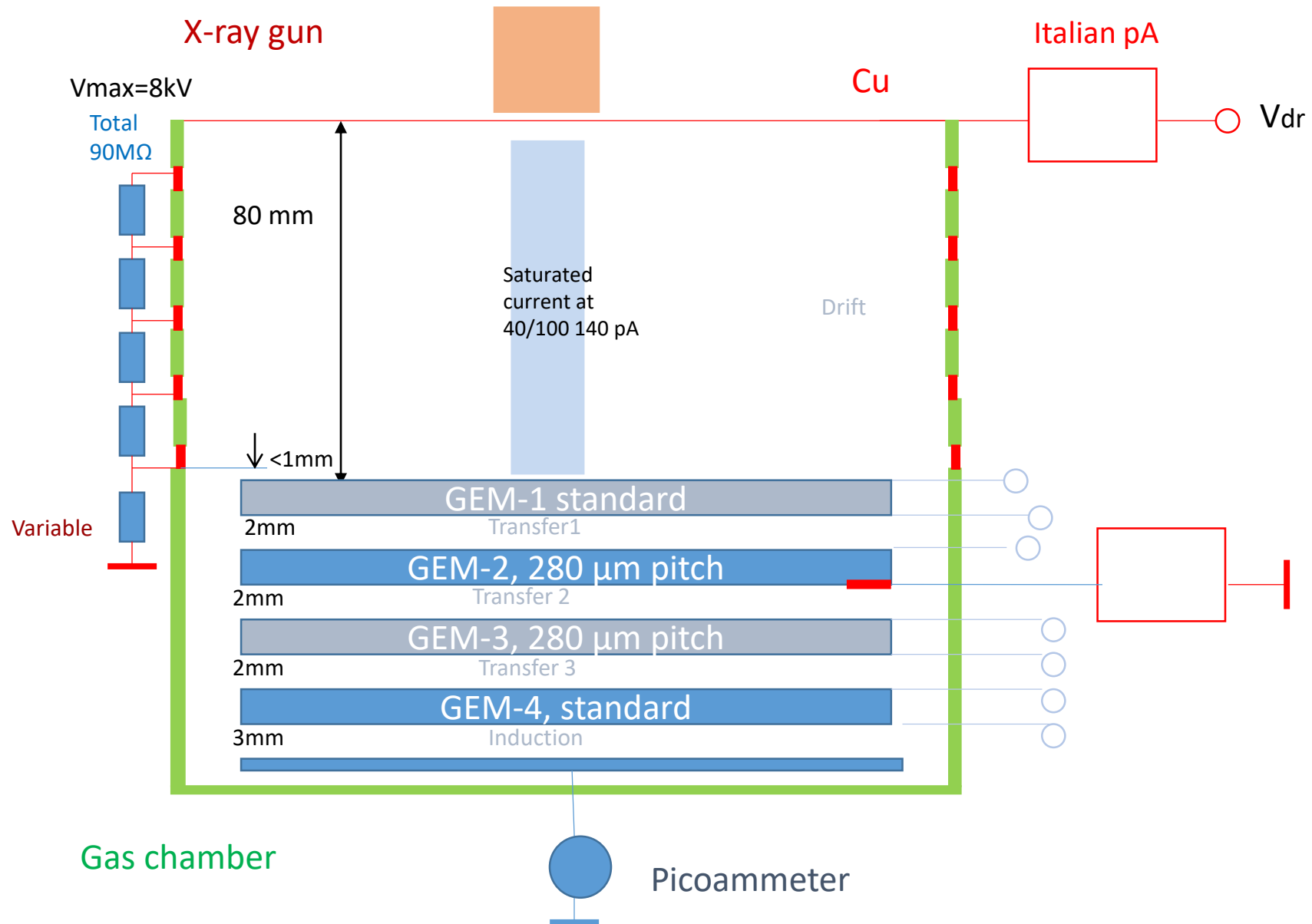


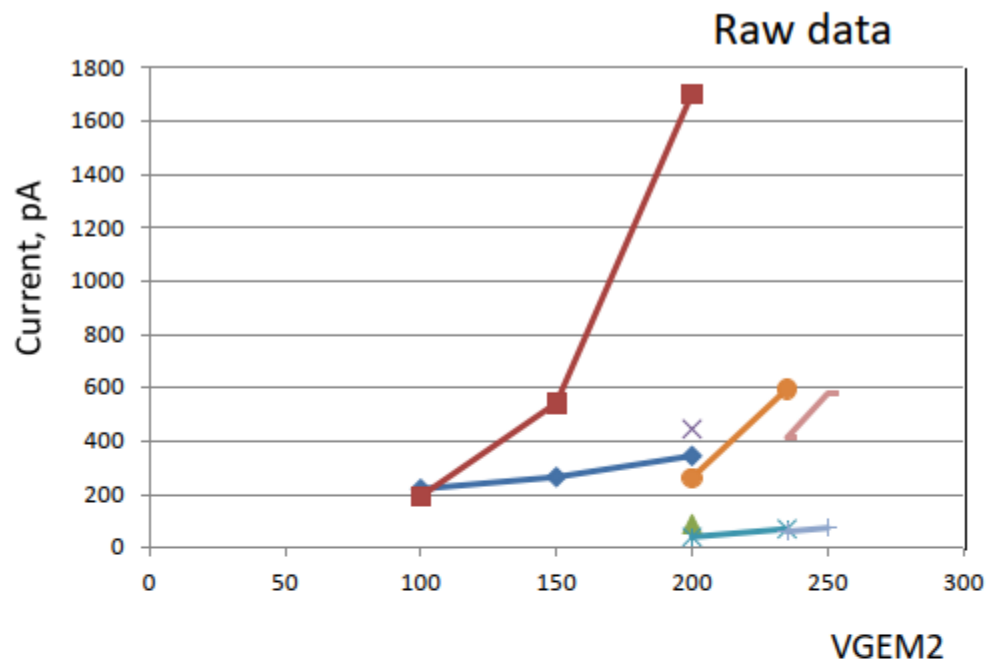
$\Delta VGEM1 = \text{const} = 245V$

At VGEM1 = 225 and Etr1 = 4 kV/cm extraction 0.47

In this case there is a strong effect of electron capture, but in ionization chamber mode negative ions are “invisible”, so results for both gas mixtures were qualitatively similar

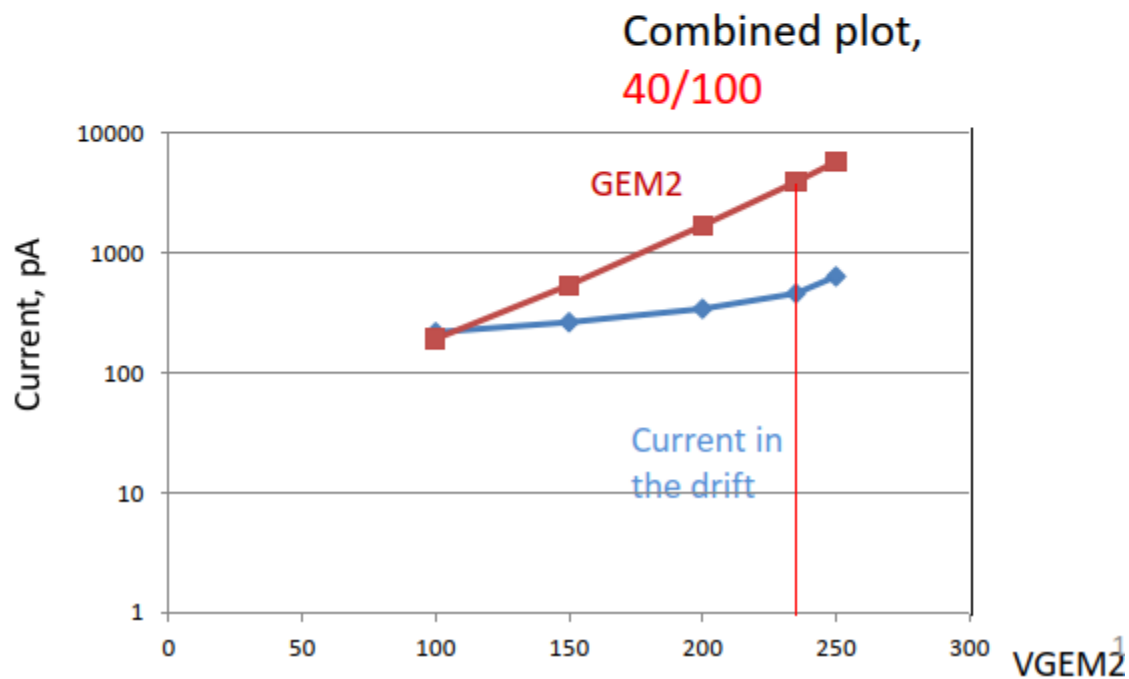
Measurements of multiplication in second GEM



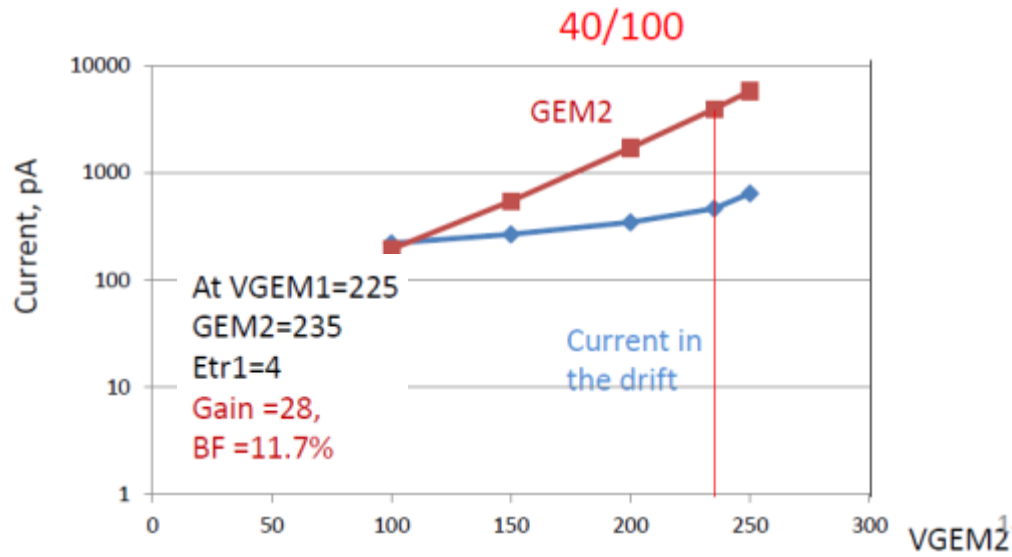


Setting:
GEM1=225,
Ert=4

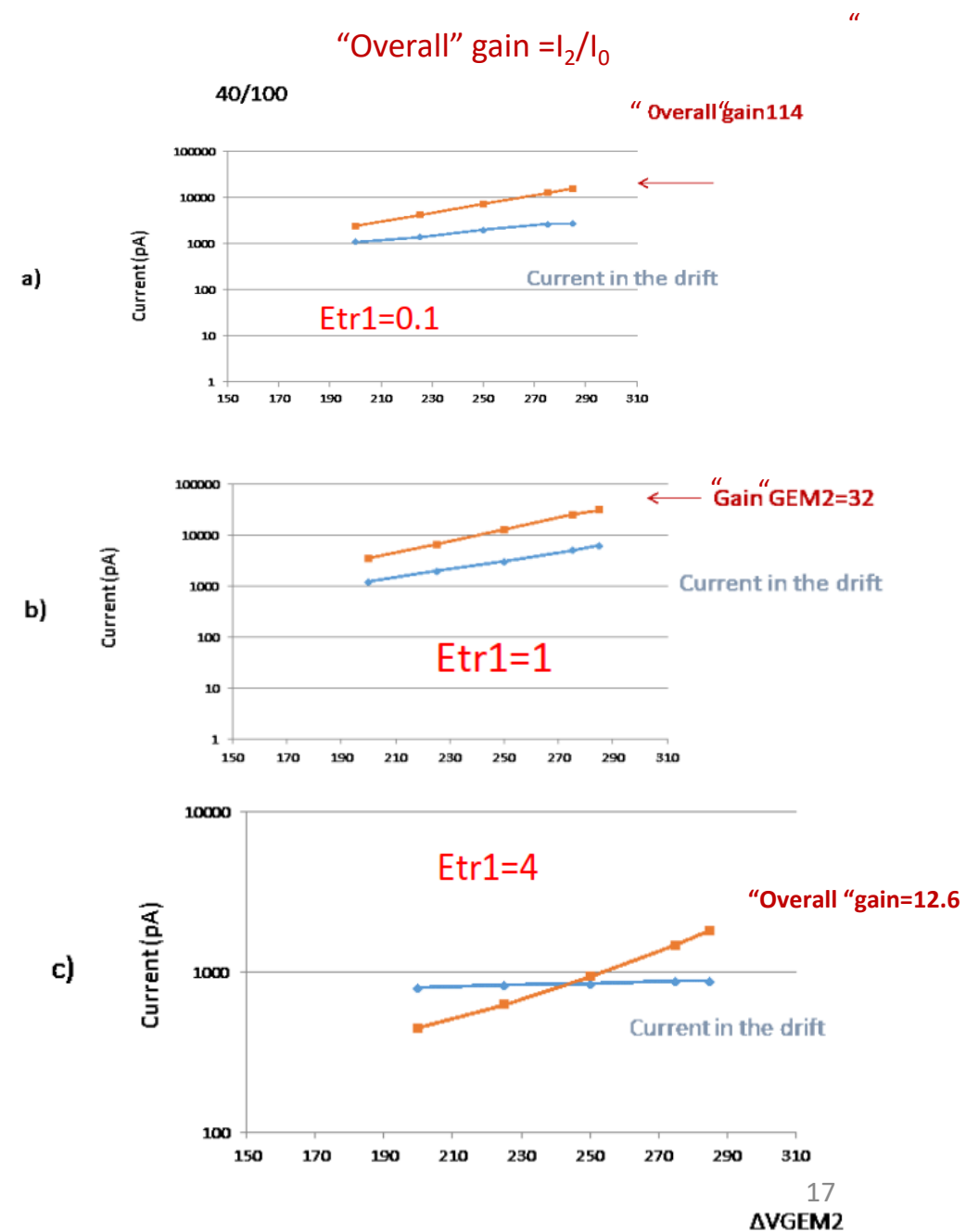
At VGEM1=225
GEM2=235
Etr1=4
Gain =28,
BF =11.7%



Ne+CO2+N2



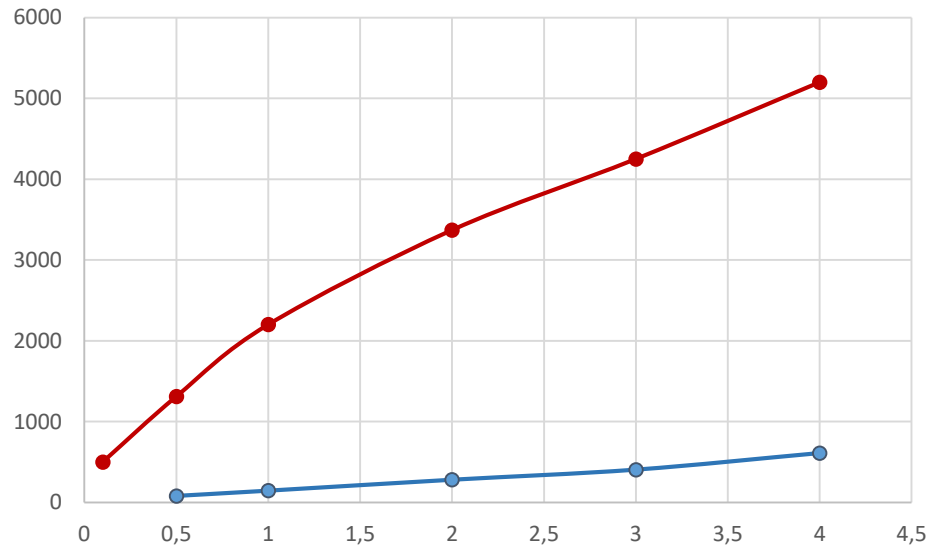
Ne+CF4



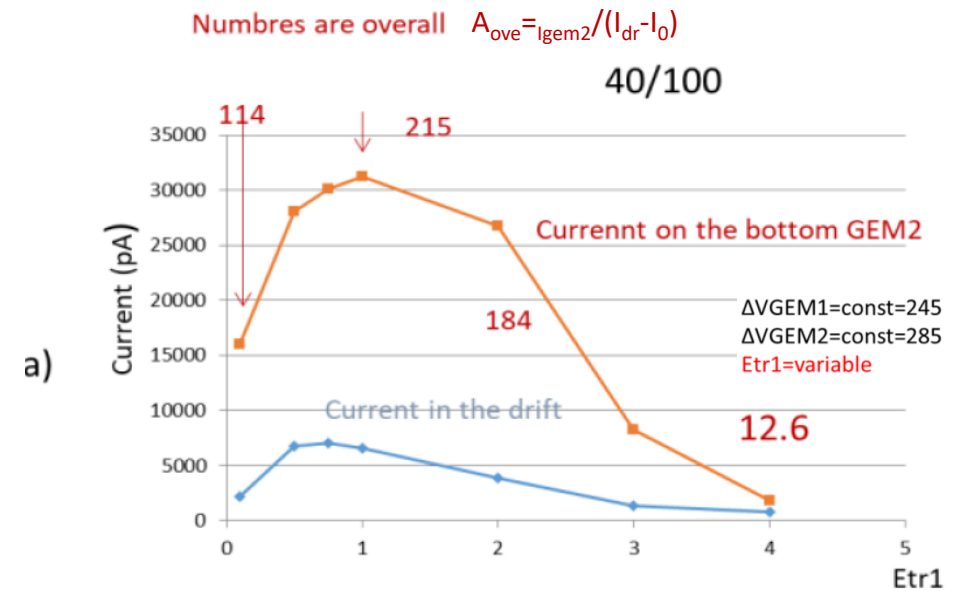
Suddenly one can see a big difference: in Ne+CF4 there is no monotonic dependence of GEMbot current as a function of the transfer field applied to its top

The same data, but presented in another variable: Etr1

Ne+CO2+N2



Ne+CF4



In the case of electronegative gases, the current generated in GEM2:

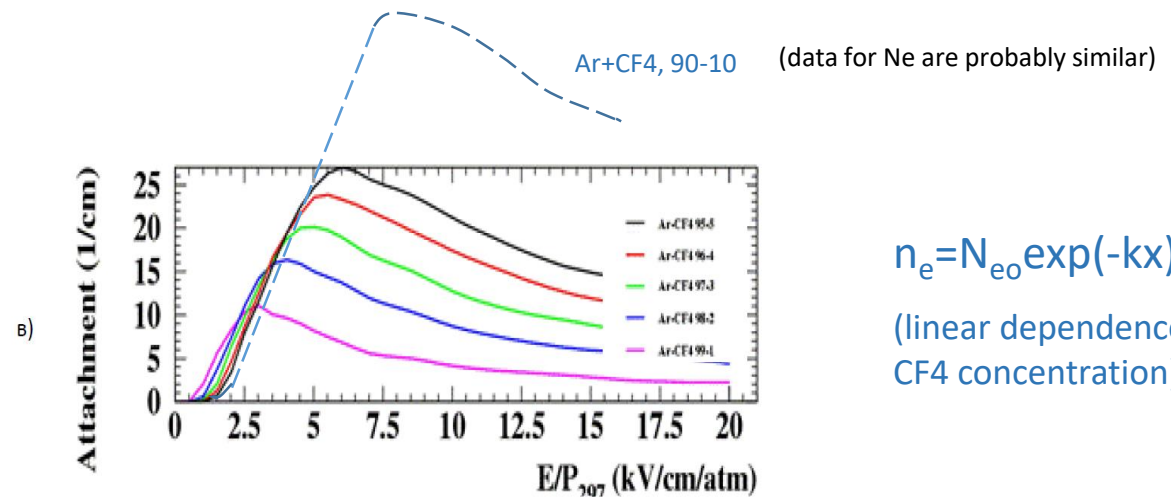
$$I_{gem2} = A_{gem2} n_e$$

$$I_{gem2} = A_{gem2} \{ I_e + I_{ni} B_{coll} (Etr_1, A) K_{da} (A) \},$$

where B_{coll} is “collection efficiency” - what proportion of negative ions falls into the holes of the lower GEM, and $K_{da} (A)$ is the coefficient indicating how much electronegative ions decay in the holes of the GEM (the so-called “disattachment process,” supposedly due to impact ionization), freeing free electrons, then initiating avalanches.

Therefore the possible qualitative explanations are

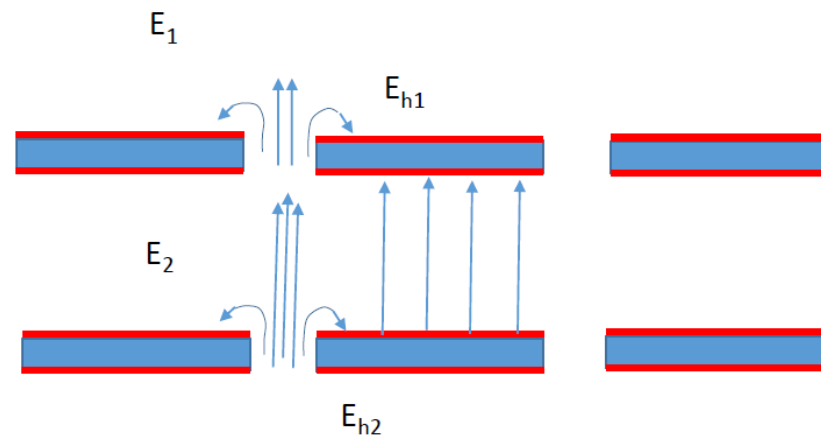
Capture of electron
(space charge can change current flows)



$$n_e = N_{e0} \exp(-kx)$$

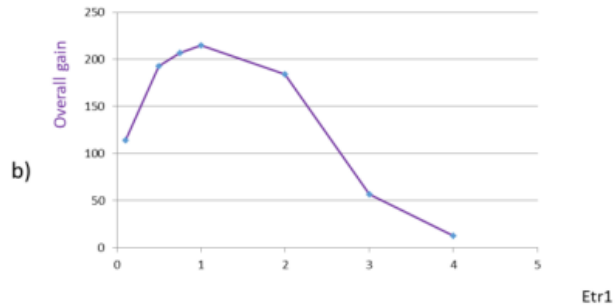
(linear dependence from CF4 concentration)

Collection between hole
(effect well known for positive ions)



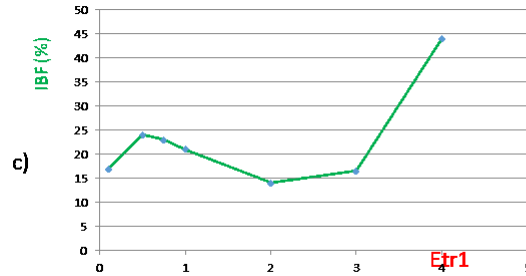
All dependence become then exotics

“Visible” overall gain



Note: gain in GEM2 = const!

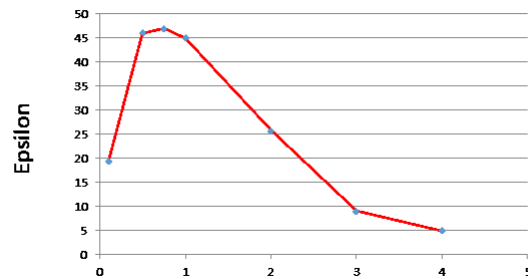
Current:



$\Delta VGEM1 = \text{const} = 245$
 $\Delta VGEM2 = \text{const} = 285$
 Etr1 = variable

d)

$$\epsilon_{IBF} = A n_{IBF} - 1$$

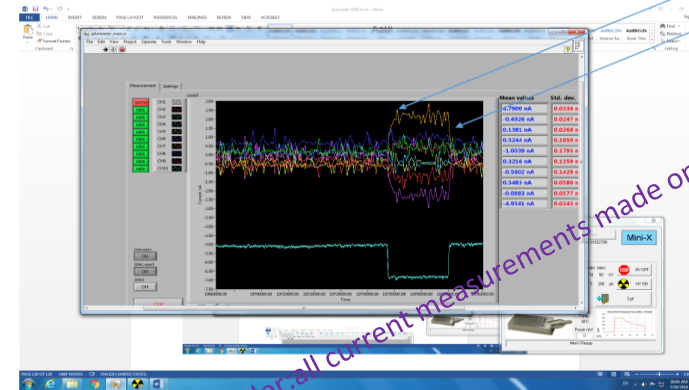


Possible reasons:

I_{ni} changes due to the space charge
 (but in our case one can neglect this effect),
 B_{coll} changes due to the low ion diffusion
 (I_e is negligible at high E)

The first was checked experimentally

Example of histograms of current from GEM electrodes, when the X-ray gun was on and off



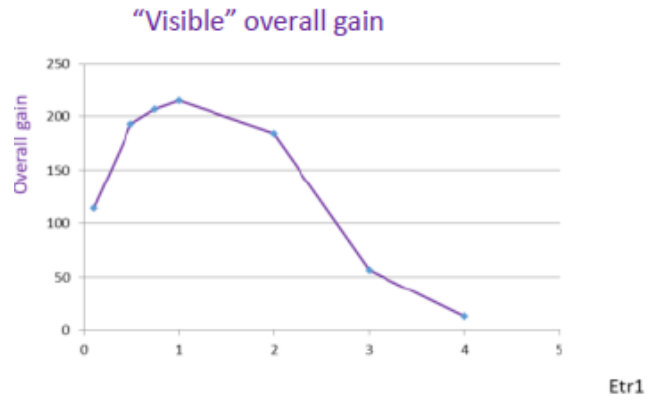
- D=1500
- T1=100
- B1=1000
- T2=900
- B2=600
- T3=500
- B3=200
- T4=100
- B4=0

Reminder: all current measurements made one of us in WIS

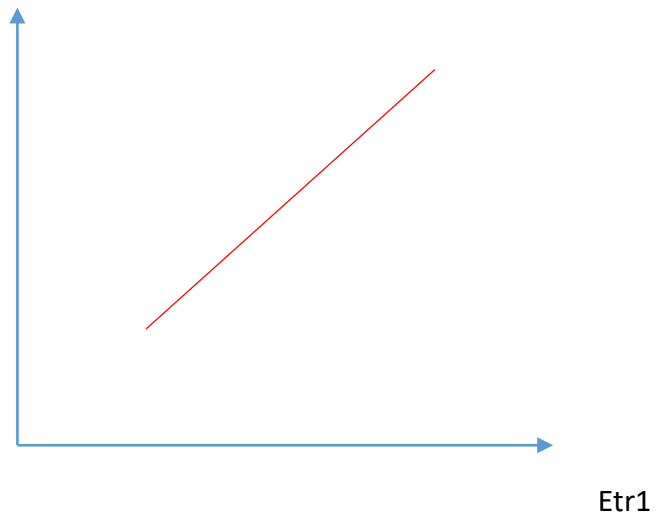
The second effect is well known

In any case, simulations are needed to clarify the reasons

Ions generated in holes



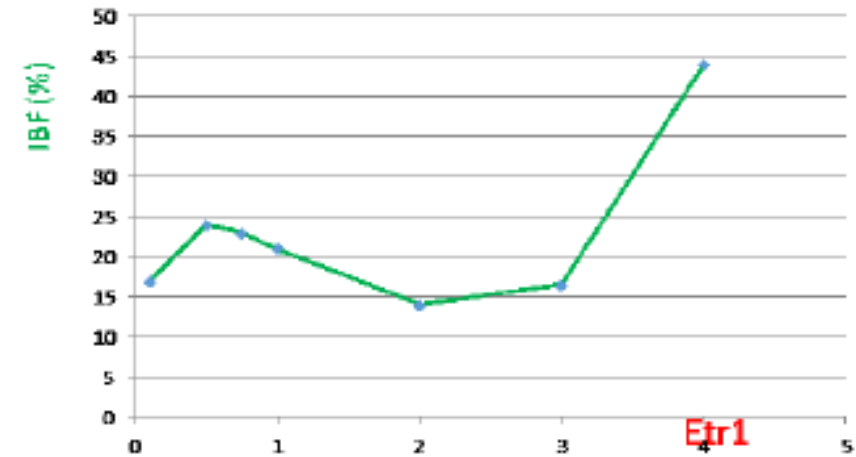
Extracted positive ions



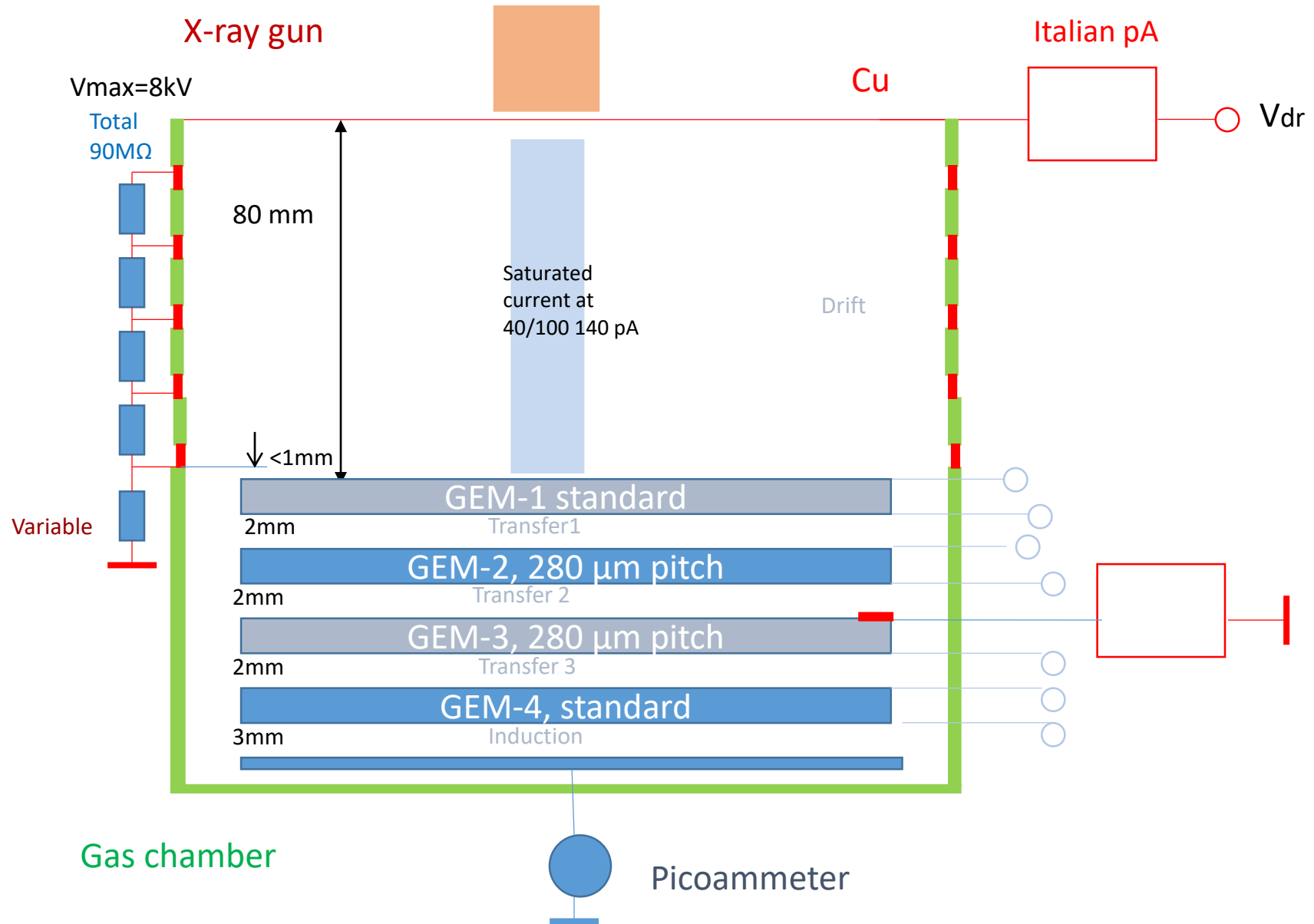
Positive ions backflow:

$$\sim A_{\text{Vis. overall}} X_{\text{extr}}$$

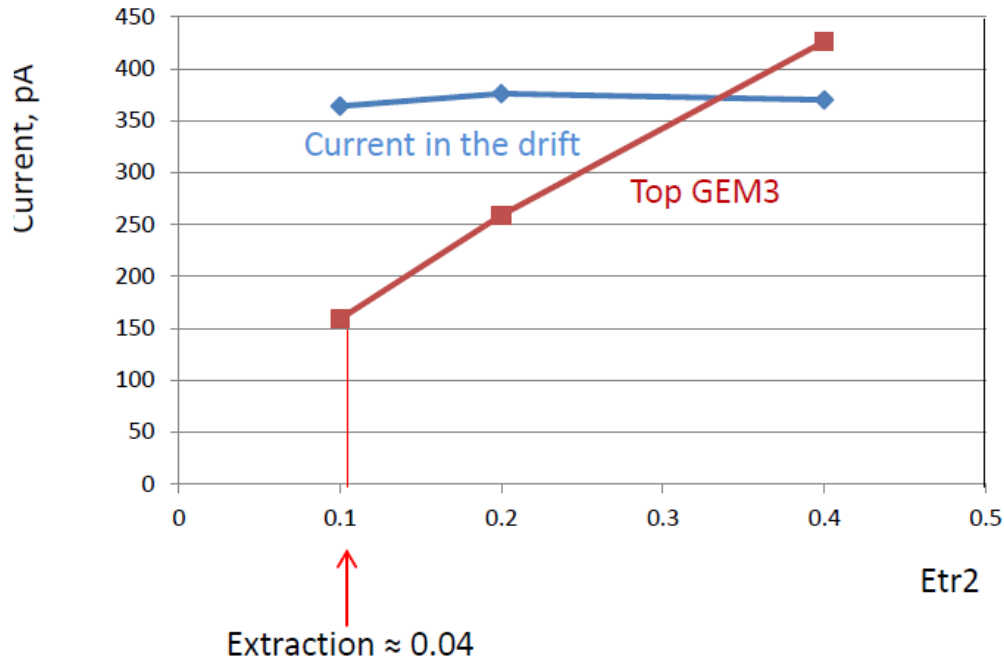
Experimental data for IBF



Measurements of extraction from second GEM



Ne+CO2+N2

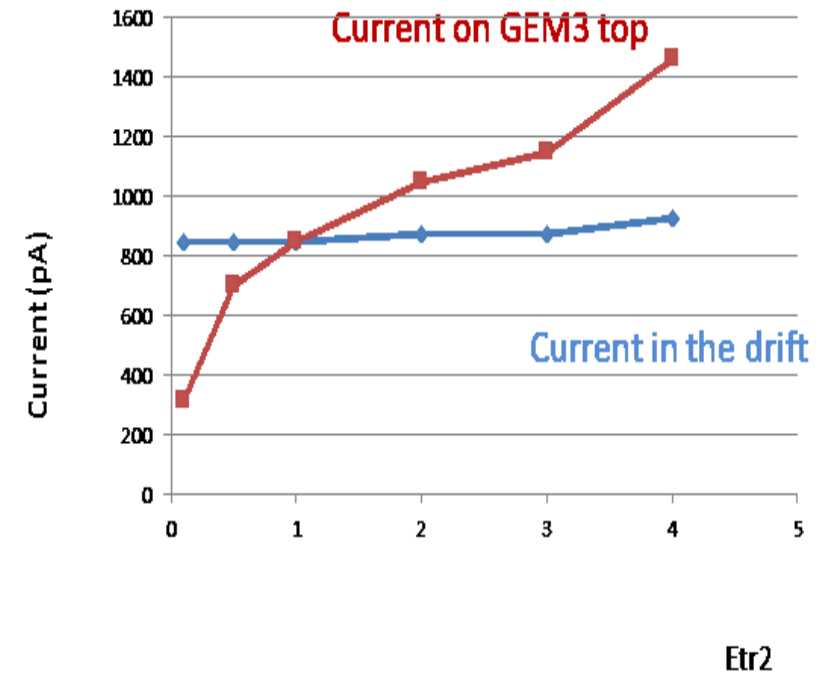


Reiner data

Ne+CF4

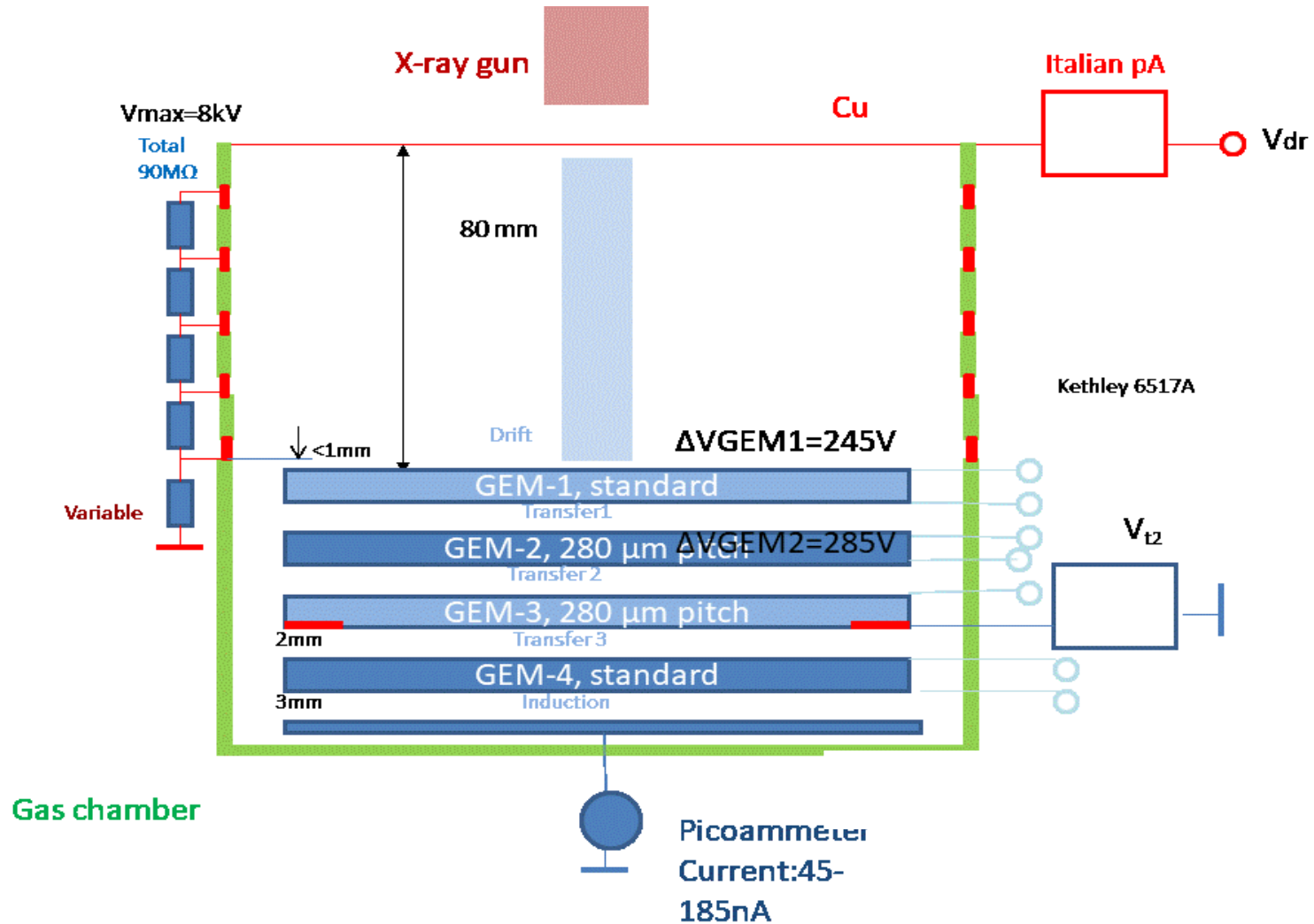
$\Delta VGEM1=245$
 $\Delta VGEM2=285$
Etr1=4
Etr2=variable

40/100

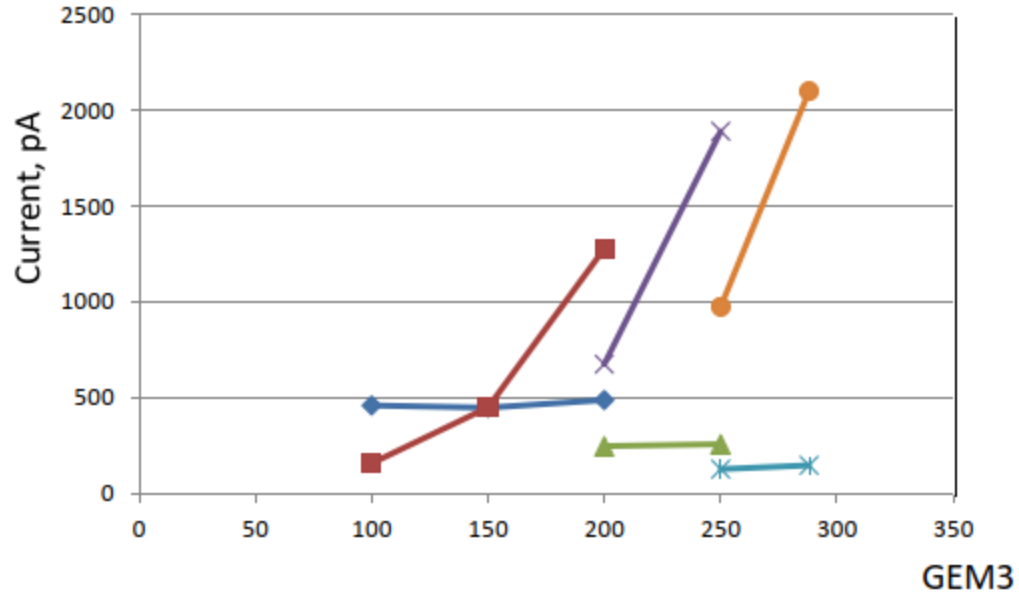


...again, in ionization chamber mode negative ions are “invisible”, so results are qualitatively similar

Measurements of multiplication in third GEM

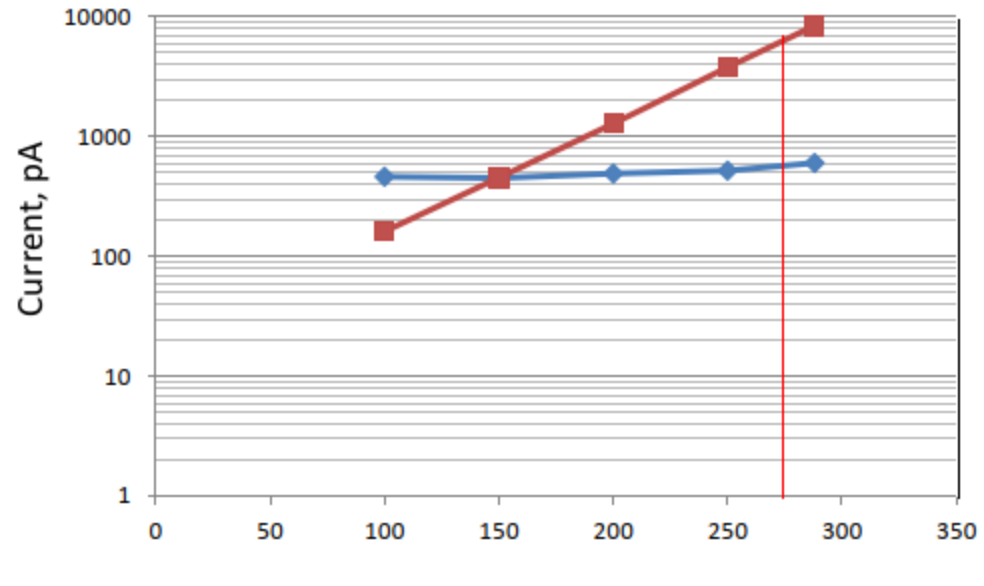


Raw data (various X-ray currents)

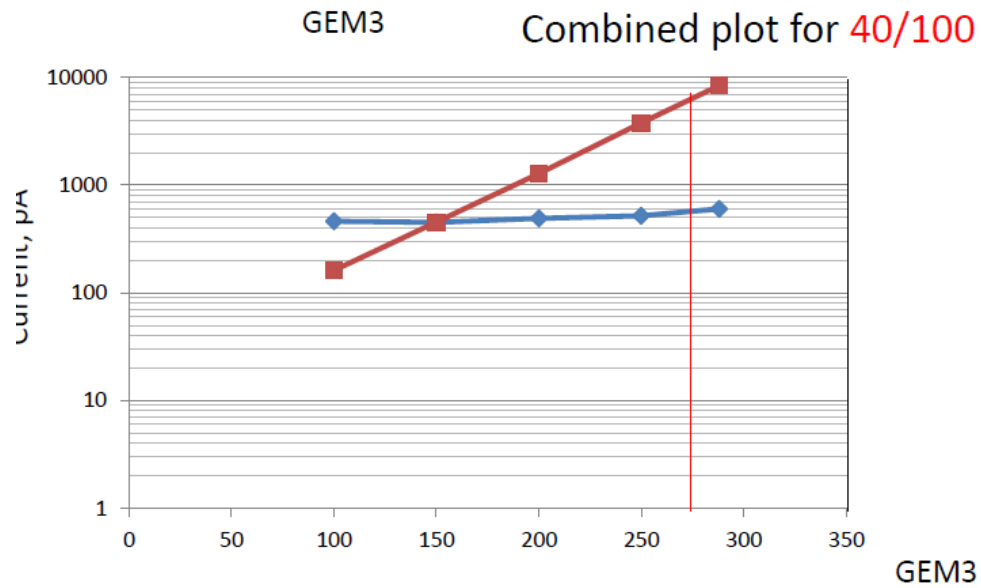


At VGEM1=225
 GEM2=235
 GEM3=272
 Etr1=4
 Et2=0.1
 Gain =42,
 BF =1.2%

Combined plot for 40/100

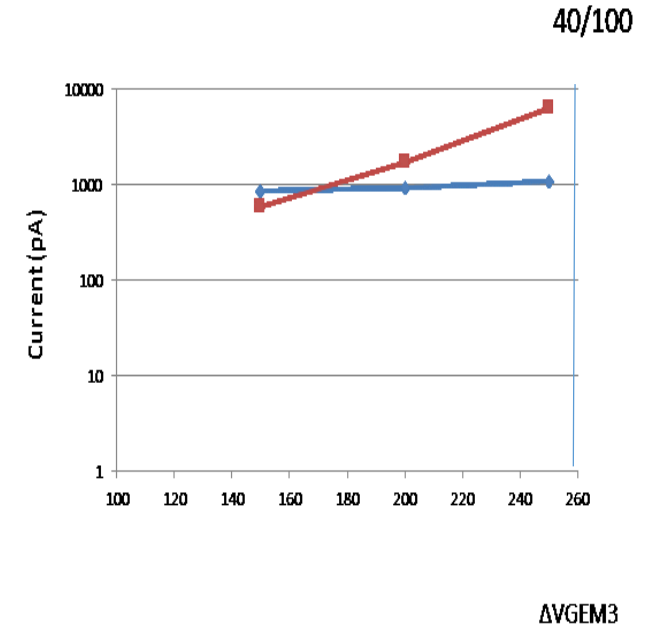


Ne+CO2+N2



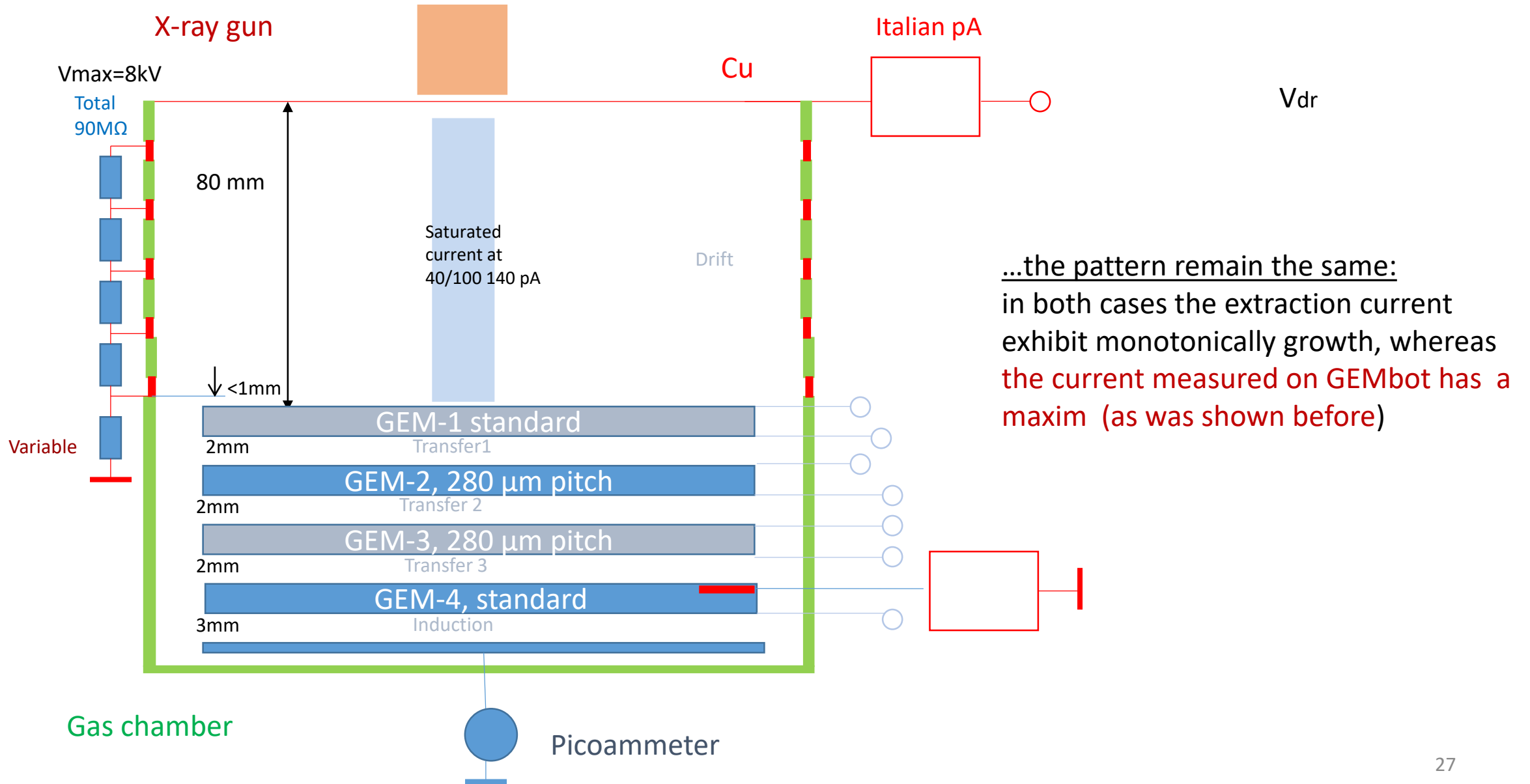
Ne+CF4

$\Delta VGEM1=245$
 $\Delta VGEM2=285$
 $\Delta VGEM3=variable$
 $E_{tr1}=4$
 $E_{tr2}=0.75$



...as expected-exponential multiplication, so no surprises

Measurements of extraction from third GEM and from fourth GEM

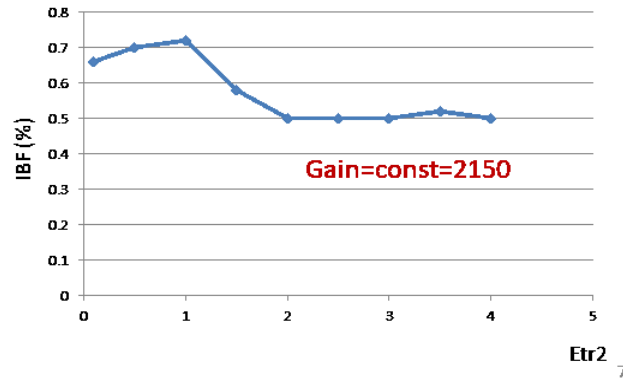
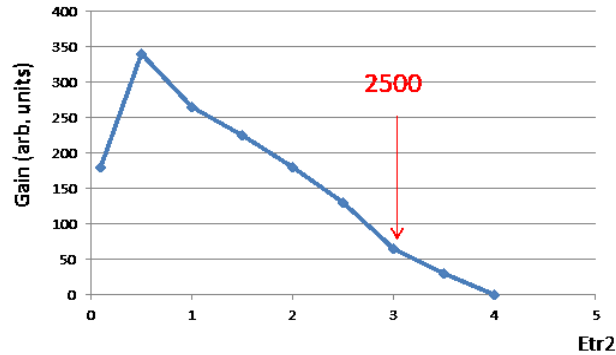


Ne+CF4

Were there any radical changes when all voltages were applied to the quadrupole GEM ?

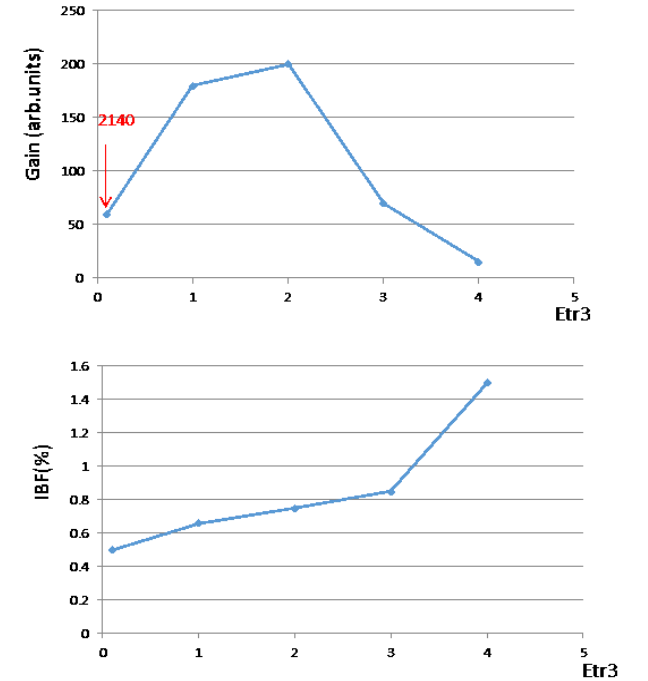
Ftr2 scan

$\Delta V_{GEM1}=245$
 $\Delta V_{GEM2}=285$
 $\Delta V_{GEM3}=320$
 $\Delta V_{GEM4}=360$
 $E_{tr1}=4$
 $E_{tr2}=0.1-4$
 $E_{tr3}=0.1$
 $E_{ind}=4$



$\Delta V_{GEM1}=245$
 $\Delta V_{GEM2}=285$
 $\Delta V_{GEM3}=320$
 $\Delta V_{GEM4}=360$
 $E_{tr1}=4$
 $E_{tr2}=3$
 $E_{tr3}=0.1-4$
 $E_{ind}=4$

... The same behaviour...

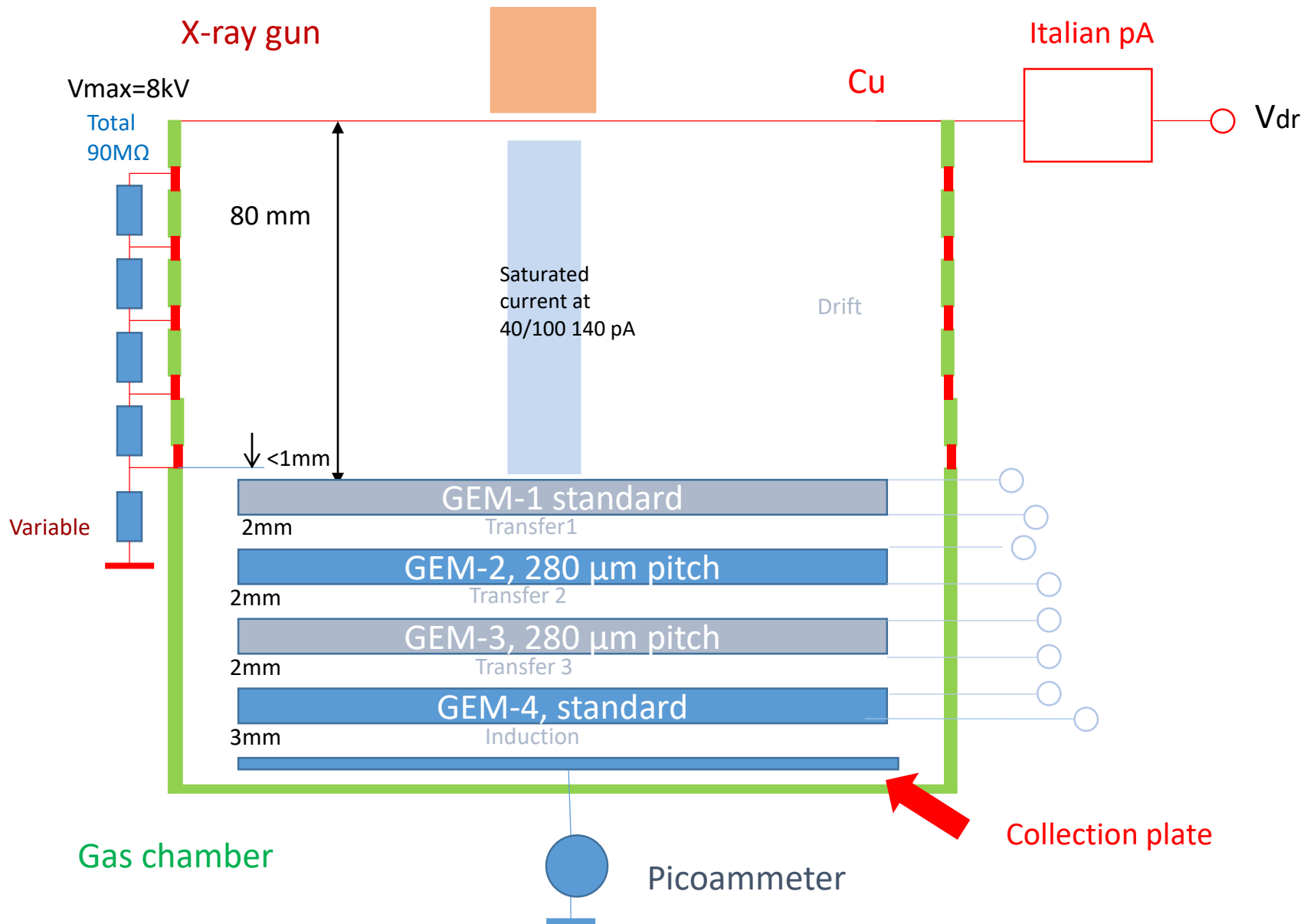


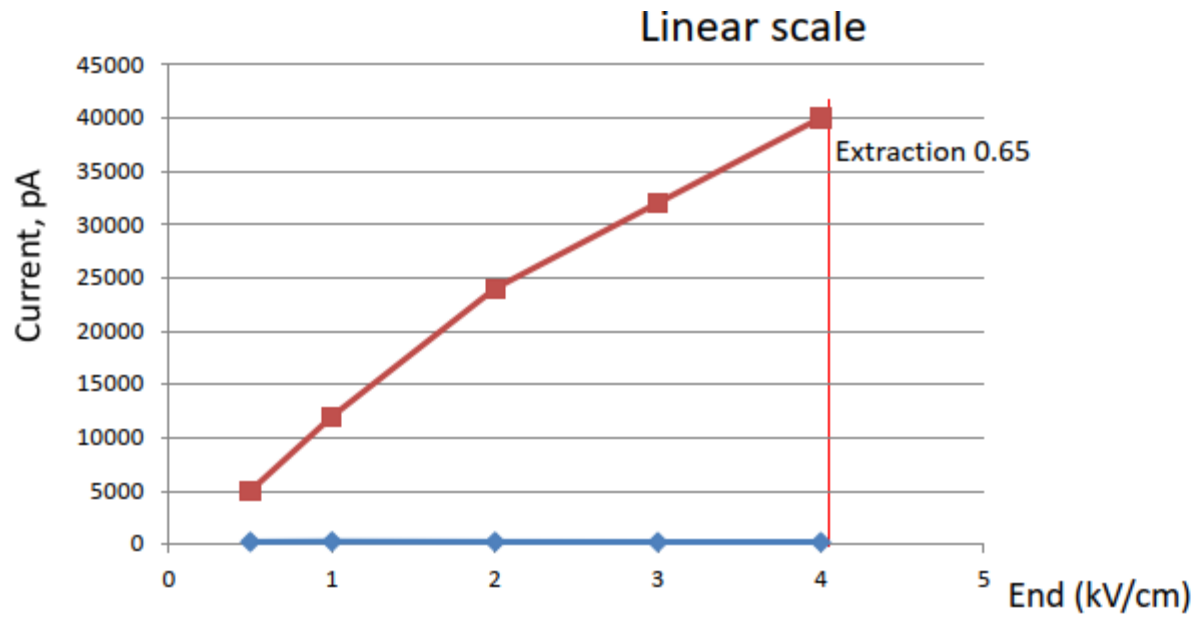
GEM3bot

GEM4bot

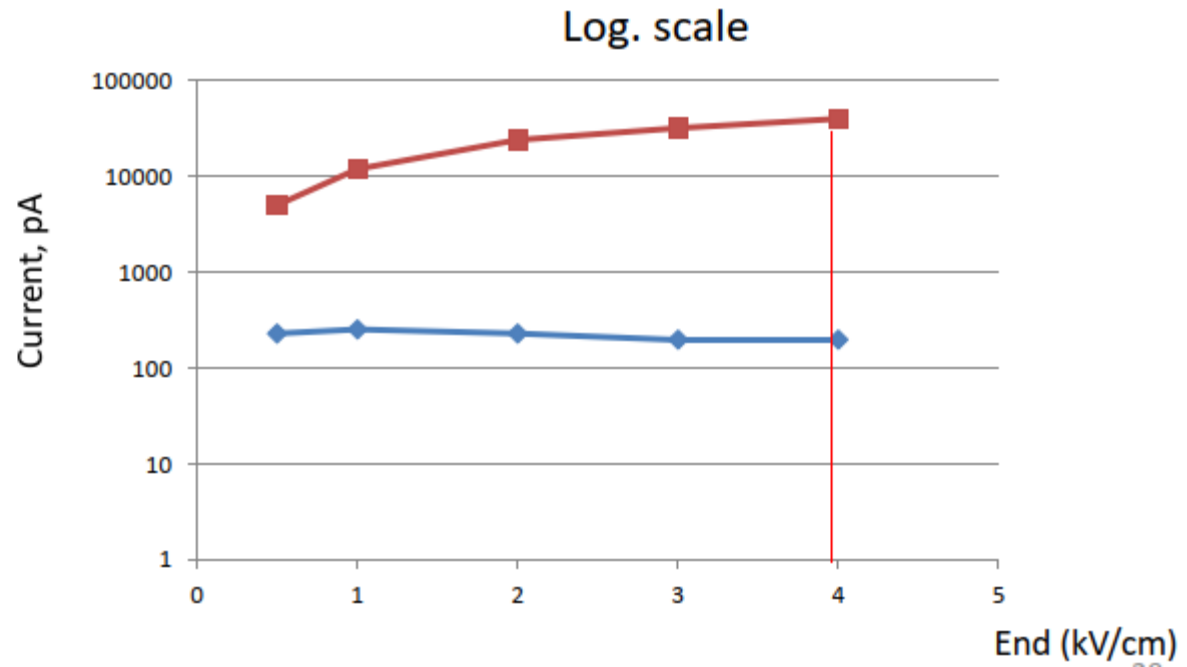
...actually no..

Measurements at the collection plate

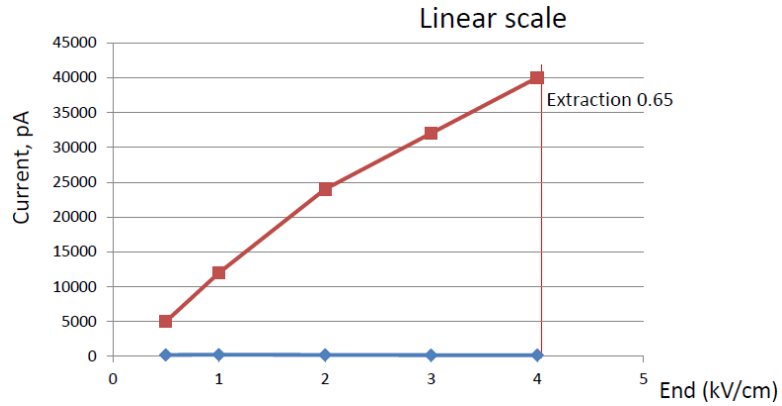




40/20



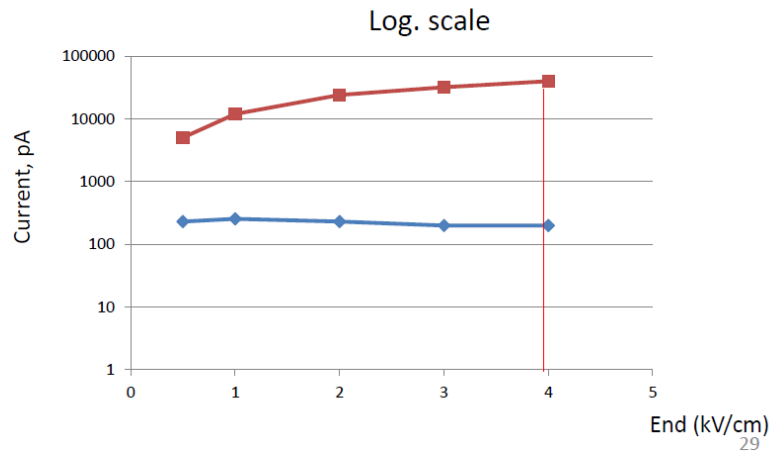
Ne+CO2+N2



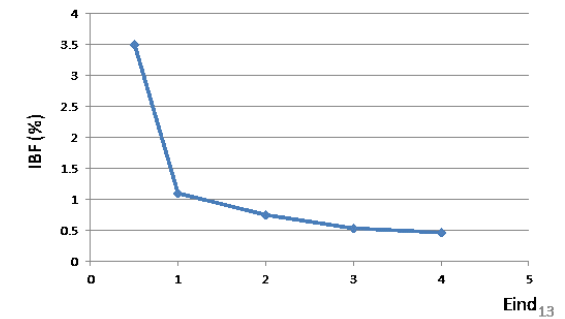
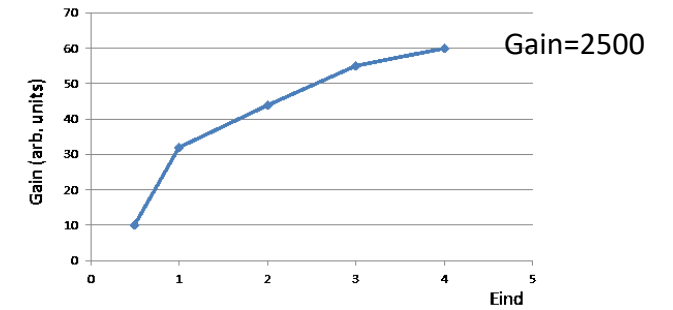
40/20

Ein scan

$\Delta VGEM1=245$
 $\Delta VGEM2=285$
 $\Delta VGEM3=320$
 $\Delta VGEM4=360$
 $E_{tr1}=4$
 $E_{tr2}=3$
 $E_{tr3}=0.1$
 $E_{ind}=0.5-4$



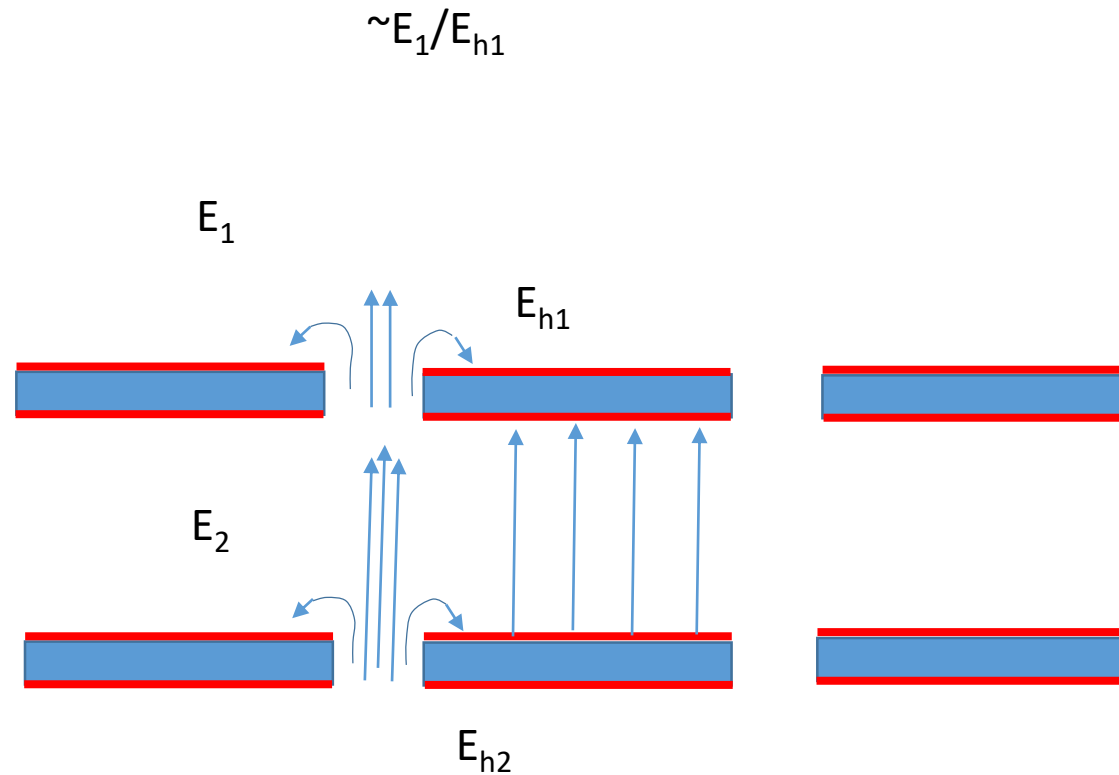
Ne+CF4



Qualitatively results are similar. In both cases the current was generated by negative carries: electrons and negative ions

Voltage setting optimization

Positive ions motion



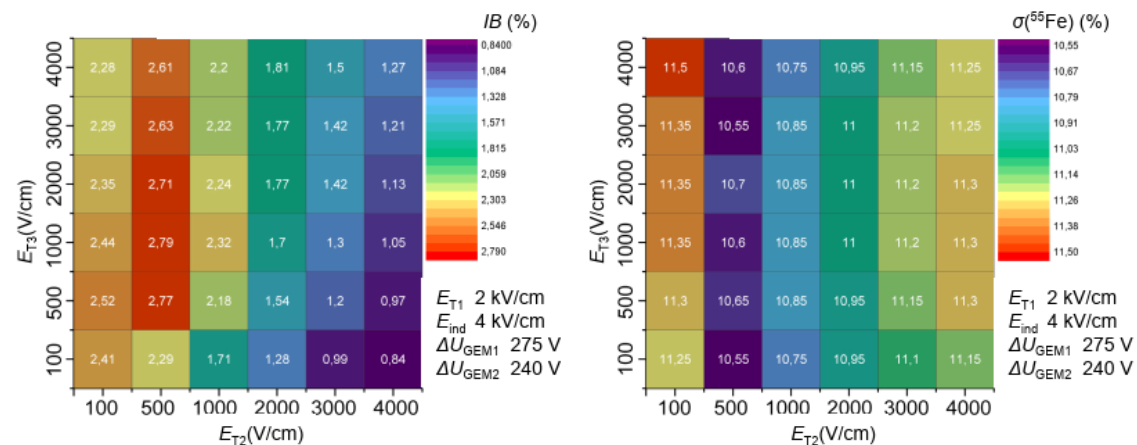
If one ignore a space charge effect
(which is valid at ALICE condition)
negative ions do not affect positive
ion movement

Therefore, roughly speaking,
one can search for the optimum
voltage settings around one found
For Ne+CO₂+N₂

One of the best setting tested so far

$\Delta V_{GEM1}=245$
 $\Delta V_{GEM2}=285$
 $\Delta V_{GEM3}=320$
 $\Delta V_{GEM4}=360$
 $E_{tr1}=4$
 $E_{tr2}=3$
 $E_{tr3}=0.1$
 $E_{ind}=4$

IBF=0.5 %
At a gain *2500* -
old calibration



Example of systematic scans performed by the ALICE TPC upgrade team

2

...so, in principle, the IBF achieved in Ne+CF₄ gas mixture could as low as in Ne+CO₂+N₂. Of course, careful scans should be done for the true voltages optimization

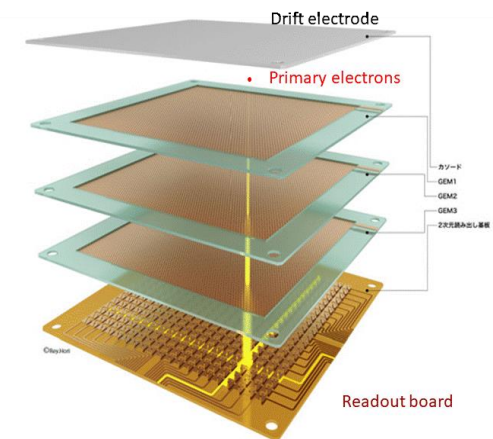


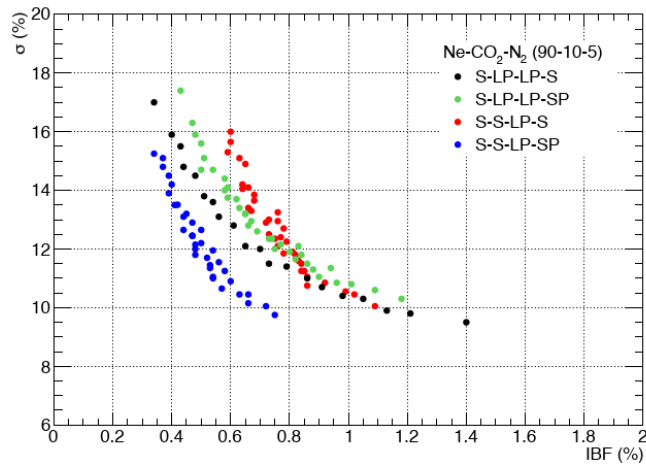
Disclaimer

The aim of the talk was not to present any systematic studies or careful voltage optimization, but rather focus on a not monotonic dependence of the gain vs. transfer field in Ne+CF₄ gas mixture

Conclusions

- “Visible “gain” of GEM in the tested Ne+CF₄ gas mixtures has a maximum as a function of the transfer field applied to its top
- This should be taken into account in a quadrupole GEM voltages optimization
- At some particular voltage setting the IBF achieved in Ne+CF₄ gas mixture was as low in the “standard” mixture Ne+CO₂+N₂ ($\approx 0.5\%$)
- However, careful scans of various parameters vs. applied voltages is missing, so the results are very preliminary and more studies should be done to prove that Ne+CF₄ gas mixture could be an alternative to Ne+CO₂+N₂

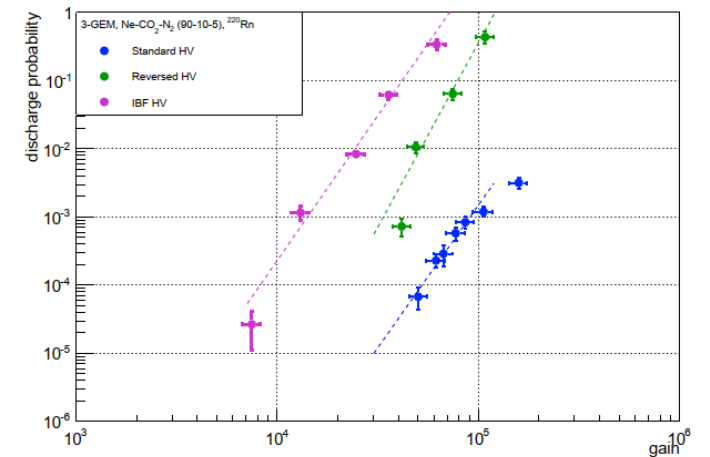




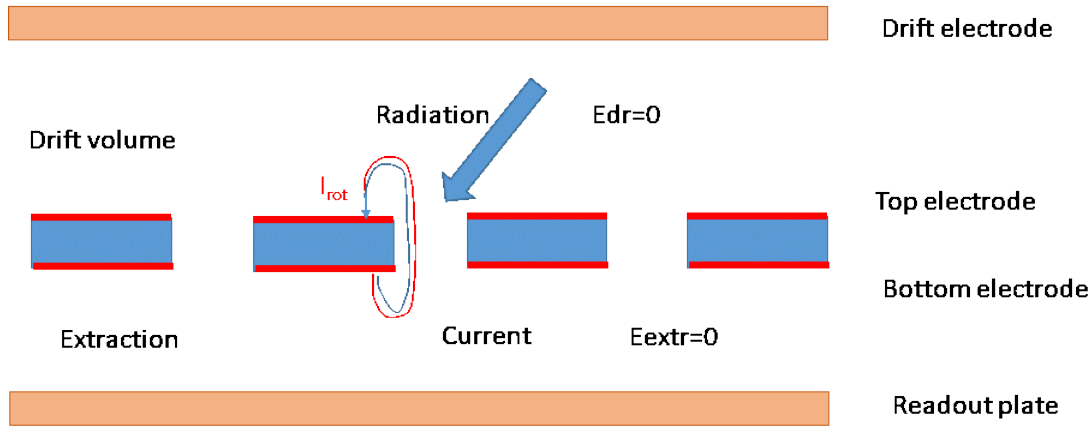
What is missing in particular?

(if one wants to go ahead with this mixture)

- Energy resolution vs. IBF measurements
 - Spark probability measurements and voltage setting optimising ensuring acceptable trade between energy resolution and spark probability
 - Long-term stability studies
 - Aging (RPC experience with water vapours))
 - Beam tests
- and, probably, much more...



Back up slides



Some basic of electrons and ions flow in GEM:

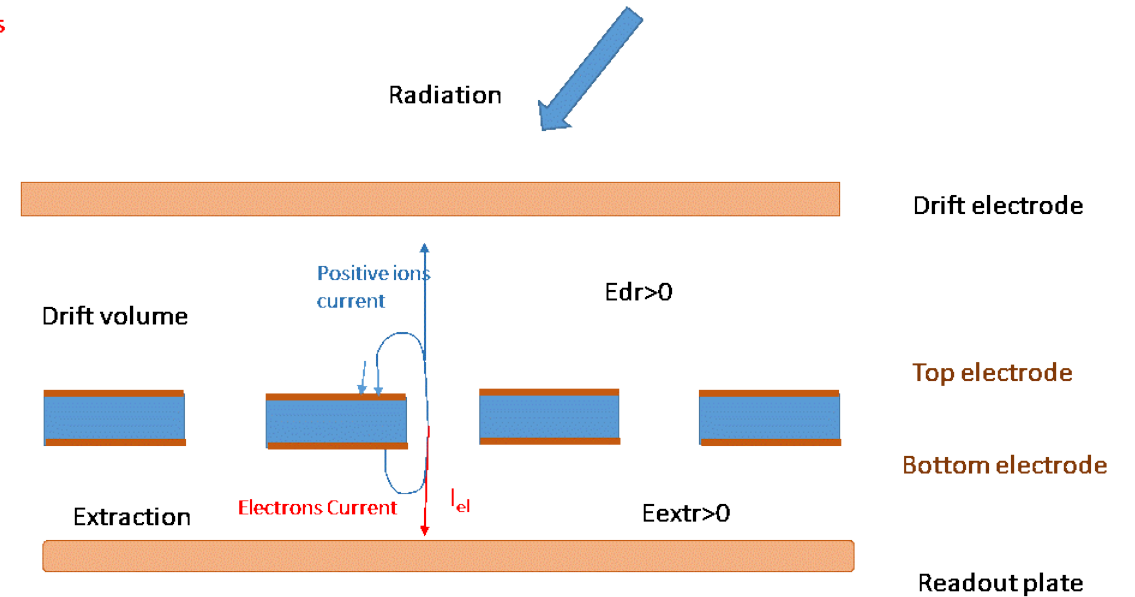
qualitative picture is quite clear, but an analytical model is quite complicated as well as simulations

In the case of E_{dr} and $E_{extr}=0$ the current I_{rot} circulate between the top and the bottom GEM electrodes

$$I_{ion} = I_{rot} f_{ion}(E_{dr})$$

$$I_{el} = I_{rot} f_{el}(E_{extr})$$

$$I_{bot1} = I_{top1} + I_{dr}$$



Similarly, if $|E_{extr}| > 0$, the extracted current (electrons) $I_{el} = I_{rot} \cdot f_{el}(E_{extr})$ where $(f_{el}(E_{extr}) < 1)$

Ne+CO2+N2

GEM1=225

GEM2=235

GEM3=272

GEM4=340

Etr1=4

Etr2=0.1

Etr3=4

Eind=4

Gain= 1800

IBF=0.5

GEM1=270

GEM2=250

GEM3=270

GEM4=340

Etr1=4

Etr2=2

Etr3=0.1

Eind=4

Gain= 2000

IBF=0.7