Lessons on Gas Detectors

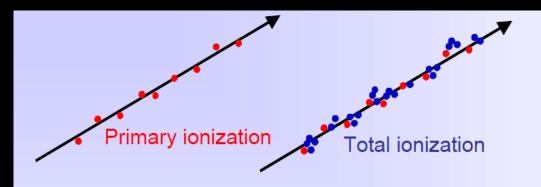
Course's aim

- Physics, architecture and construction techniques of <u>Gaseous</u> <u>Detectors</u>
- Part I (Prof. F. Tessarotto)
 - Theoretical introduction (ionization, transport, multiplication, signal formation)
 - MPGDs in general (historical introduction, MSGCs, GEMs, Micromegas and other architectures)
 - GEMs, THGEMs and Micromegas (manufacturing, performance, applications)
 - PID (introduction, RICH principle, radiators, optics, photon detectors, example of RICHs)
 - Gaseous photon detectors (general characteristics, THGEMs + Csl, perspectives)

- Part II (Prof. Verwilligen)
- Simulation activity
- The students will face a few simulation aspects of micro-pattern gaseous detectors. During this part, the students will calculate and visualize the electric field of a GEM detector.
- Thereafter the students will use the calculated electric field to generate the response to a muon passing through the GEM detector.
- Therefore, they will simulate the gas with MAGBOLTZ, simulate the ionization of a muon passing through the GEM with HEED, and track the drifting ionization electrons with GARFIELD.
- GARFIELD will then be used to simulate the avalanches initiated by those ionization electrons and finally the students will calculate the signal induced by these avalanches.

- Part III (A. Ranieri)
- Laboratory experiences
- It will be performed a few classical measurements (gain, rates, sensitivity, etc..) based on a small planar GEM detector.

The gas ionization



Fast charged particles ionize atoms of gas.

Often resulting primary electron will have enough kinetic energy to ionize other atoms.

$$n_{total} = \frac{\Delta E}{W_i} = \frac{\frac{dE}{dx}\Delta x}{W_i}$$

 $n_{total} \approx 3...4 \cdot n_{primary}$

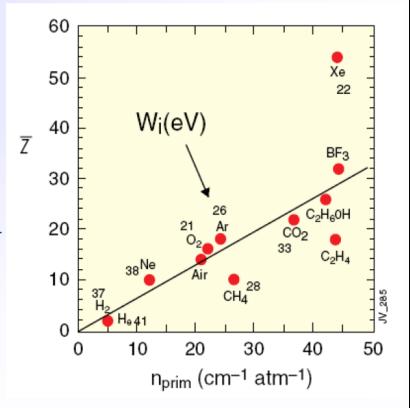
 n_{total} - number of created electron-ion pairs

 Δ_E = total energy loss

 W_i = effective <energy loss>/pair

Number of primary electron/ion pairs in frequently used gases.

Lohse and Witzeling, Instrumentation In High Energy Physics, World Scientific, 1992



The gas ionization

The actual number of primary electron/ion pairs is Poisson distributed.

$$P(m) = \frac{\overline{n}^m e^{-\overline{n}}}{m!} \qquad \qquad -\frac{L}{n} = LN\sigma_i$$

The detection efficiency is therefore limited to:

$$\varepsilon_{\text{det}} = 1 - P(0) = 1 - e^{-\overline{n}}$$

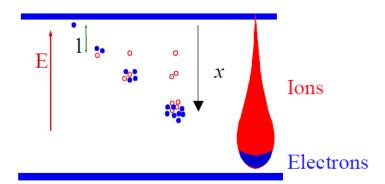
For thin layers ε_{det} can be significantly lower than 1.

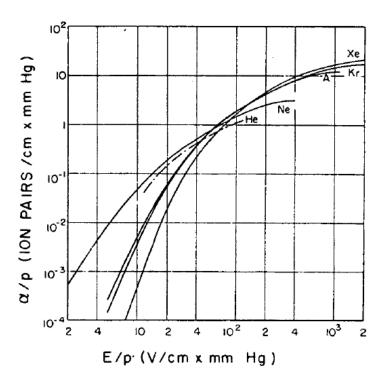
For example for 1 mm layer of Ar $n_{primary}$ = 2.5 $\rightarrow \varepsilon_{det}$ = 0.92 .

100 electron/ion pairs created during ionization process is not easy to detect.

Typical noise of the amplifier $\approx 1000~e^{-}$ (ENC) \rightarrow gas amplification .

Charge moltiplication process in uniform field





Mean free path for ionization:

$$\lambda = \frac{1}{N\sigma}$$
 N: molecules/cm³

Townsend coefficient:

$$\alpha = \frac{1}{\lambda}$$
 Ionizing collisions/cm $\frac{\alpha}{P} = f\left(\frac{E}{P}\right)$

$$\alpha = Ape^{-B\frac{P}{E}}$$

Incremental increase of the number of electrons in the avalanche:

$$dn = n \alpha dx$$

Multiplication factor (Gain):
$$M(x) = \frac{n}{n_0} = e^{\alpha x}$$

Maximum Avalanche size before discharge (Raether limit):

Raether limit: $M < 10^8 \rightarrow \alpha x < 20$

H. Raether, Electron Avalanches and Breakdown in Gases (Butterworth 1964) $\alpha(E)$ is determined by the excitation and ionization cross section of the electrons in the gas

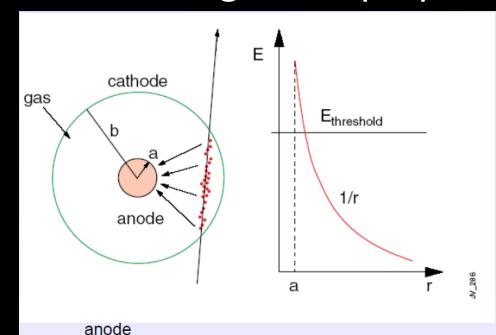
α(E) depends also on various and complex energy transfer mechanisms between gas molecules.

It has to be measured for every gas mixtures

$$M = \exp\left[\int_{x_1}^{x_2} \alpha(x) dx\right]$$

S.C. Brown, Basic Data of Plasma Physics (MIT Press, 1959)

Single wire proportional chamber



primary electron

Electrons liberated by ionization drift towards the anode wire.

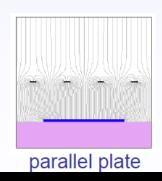
Electrical field close to the wire (typical wire \emptyset ~few tens of μ m) is sufficiently high for electrons (above 10 kV/cm) to gain enough energy to ionize further \rightarrow avalanche – exponential increase of number of electron ion pairs.

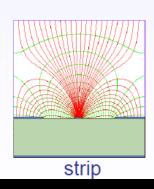
$$E(r) = \frac{CV_0}{2\pi\varepsilon_0} \cdot \frac{1}{r}$$

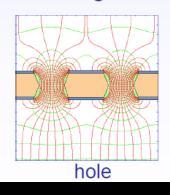
$$V(r) = \frac{CV_0}{2\pi\varepsilon_0} \cdot \ln\frac{r}{a}$$

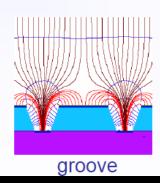
$$C - \text{capacitance/unit length}$$

Cylindrical geometry is not the only one able to generate strong electric field:



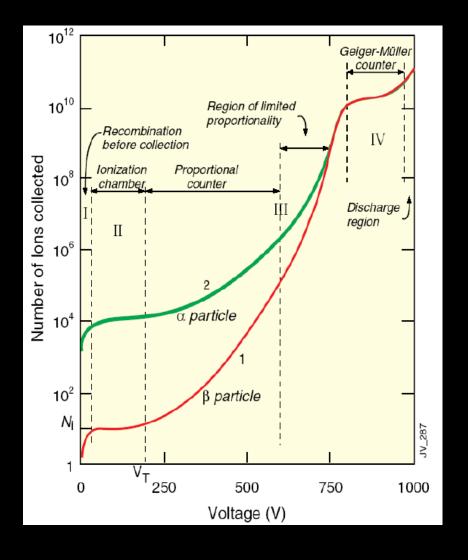






SWPC – Operation Modes

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ionization mode - full charge collection, but no
charge multiplication;
gain ~ 1
proportional mode - multiplication of ionization
starts; detected signal proportional to original
ionization → possible energy measurement (dE/dx);
secondary avalanches have to be quenched;
gain \sim 10^4 - 10^5
limited proportional mode (saturated, streamer) -
strong photoemission; secondary avalanches
merging with original avalanche; requires strong
quenchers or pulsed HV; large signals → simple
electronics;
gain \sim 10^{10}
Geiger mode – massive photoemission; full length
of the anode wire affected; discharge stopped by
HV cut; strong quenchers needed as well
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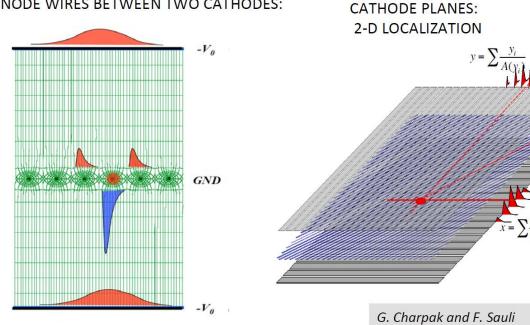
The invention of MWPC

MULTIWIRE PROPORTIONAL CHAMBER (MWPC)

TWO-TRACK RESOLUTION

CHARGE INDUCTION ON

THIN ANODE WIRES BETWEEN TWO CATHODES:



G. Charpak et al, Nucl. Instr. and Meth. 62(1968)262

Two-dimensional coordinate readout:

Nucl. Instr. and Meth. 113(1973)381

$$X = \sum \frac{X_i A_i(X)}{A(X)} \qquad Y = \sum \frac{Y_i A_i(X)}{A(Y)}$$

 X_i , Y_i : Coordinates of the strips $A_i(X)$, $A_i(Y)$: Charge on strips A(X), A(Y): Total charge

A major revolution occurred in 1968, when Charpak invented Georges the Multiwire Chamber(MWPC), a Proportional detector outperforming by orders of magnitude the rate capability of contemporary devices. Consisting of a grid of thin, parallel anode wires between two cathode planes, on application of suitable voltages the device collects and amplifies, by avalanche multiplication, the tiny ionization clusters released in a gas by Ionizing radiation, permitting detection with electronics means.

Pro's:

Space resolution improvement ($\Delta x=d/\sqrt{12}$) Good rate capability

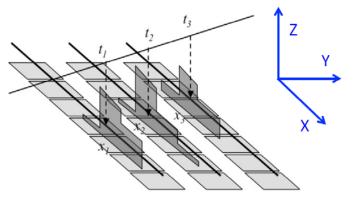
Con's:

Increasing of Electric field attraction for narrower wires distance (>> stability and mechanical issue)

Adding third dimension

TIME PROJETION CHAMBER: FULL 3-D LOCALIZATION

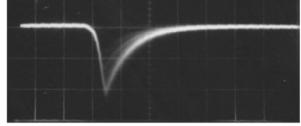
TWO-TRACK RESOLUTION



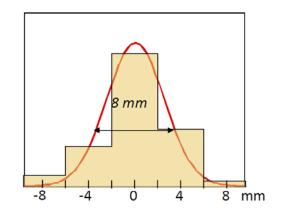
Y: ANODE WIRE Δ Y ~ 4 mm

VOLUME RESOLUTION: ΔV^{\sim} 50 mm³

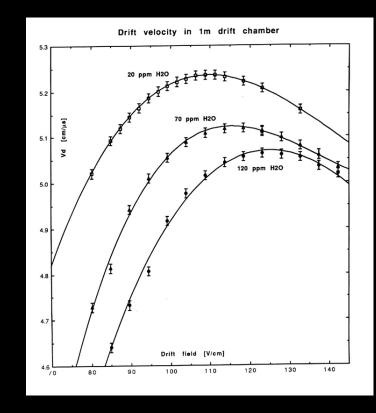
Z: DRIFT TIME Δ Z ~ 10 mm (200 ns):



X: PAD ROWS $\Delta X \sim 10 \text{ mm}$:



The MWPC was quickly adopted and successfully used by many experiments. Owing to the development work in many groups, the basic structure evolved and diversified into several more sophisticated devices exploiting the measurement of the collection or drift time of electrons to the anodes or the recording of the signals induced on cathodes to improve time and space resolutions.



Big TPC construction

D. Decamp et al, Nucl. Instr.

WO-TRACK RESOLUTION TIME PROJETION CHAMBER (TPC) FIRST TPC: PEP-4 AT SLAC (1975) D.R. Nygren and J. N. Marx, Physics Today No.31 Vol. 10(1978) ALEPH TPC AT CERN-LEP (1989) WIRE CHAMBERS

Good dE/dx resolution requires

long track length

large number of samples/track

good calibration, no noise, ...

- Energy loss (dE/dx) depends on the particle velocity (Bethe-Block)
- The mass of the particle can be identified by measuring simultaneously momentum and dE/dx (ion pairs produced)
- Major problem is the large Landau fluctuations on a single dE/dx sample

- Field cage
- Gas system
- Wire chambers
- Gating
- Magnetic field
- Laser system
- Electronics

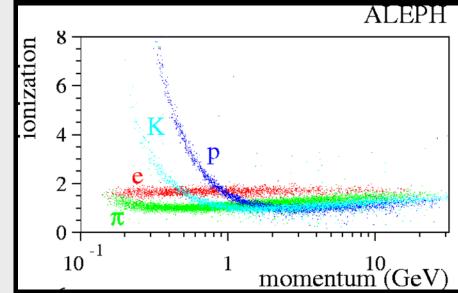
Dimensions cylinder $4.7 \times 1.8 \text{ m}$

Drift length 2x2.2 m

Electric field 110 V/cm

Electrodes

copper strips (35 μ m & 19 μ m thickness, 10.1 mm pitch, 1.5 mm gap) on Kapton



Space resolution: $\sigma_{r\phi} = 170 \mu m$ $\sigma_{z} = 700 \mu m$

Typical gas parameters & their influences on TPC performances

Typical mixtures: Ar(91%)+CH₄(9%),

 $Ar(90\%)+CH_4(5\%)+CO_2(5\%)$

Operation at atmospheric pressure

Properties:

Drift velocity (~ 5 cm/ μ s)

Gas amplification (~7000)

Signal attenuation (electron

attachment) (<1%/m)

Parameters to control and monitor:

Mixture quality (change in amplification)

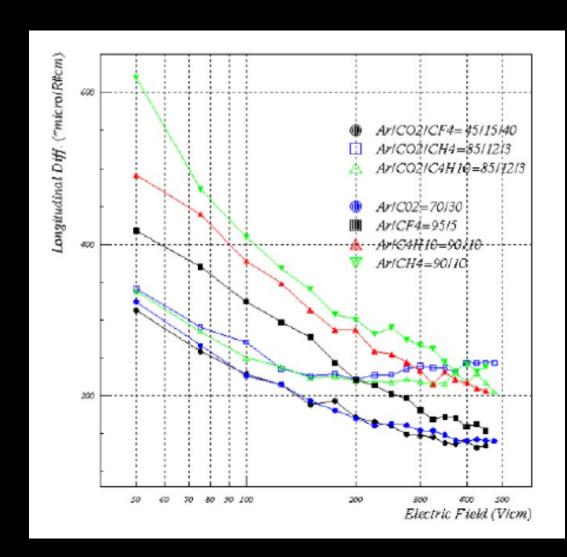
 O_2 (electron attachment, attenuation)

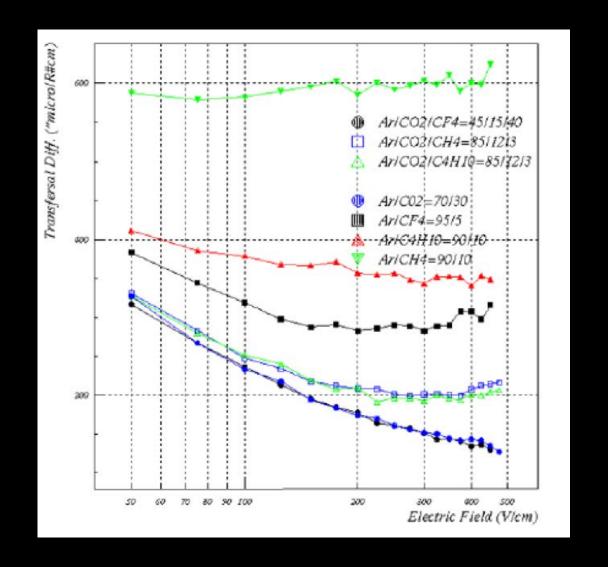
H₂O (change in drift velocity, attenuation)

Other contaminants (attenuation)

Parameter change	Drift velocity, v _d	Effect on gas amplification, A	Signal ettenuation by electron attachment
0.1% ∆CH ₄	0.4 %	-2.5% for $A = 1 \times 10^4$	
10 ppm O ₂	Negligible up to 100 ppm	Negligible up to 100 ppm	0.15%/m of drift
10 ppm H ₂ O	0.5 %	Negligible at 100 ppm	< 0.03% /m of drift
1 mbar	Negligible if at max.	-(0.5%-0.7%)	

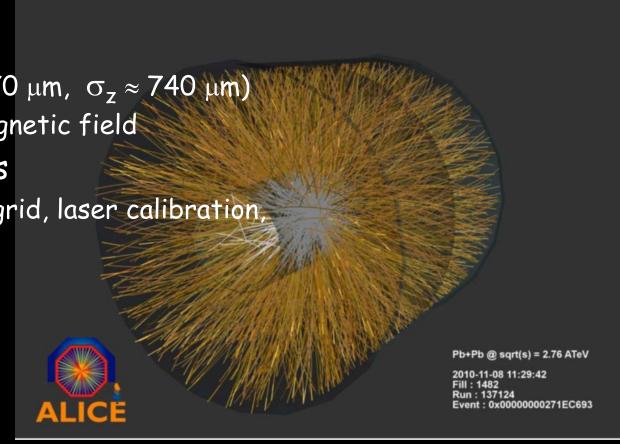
Lateral and longitudinal charge diffusion



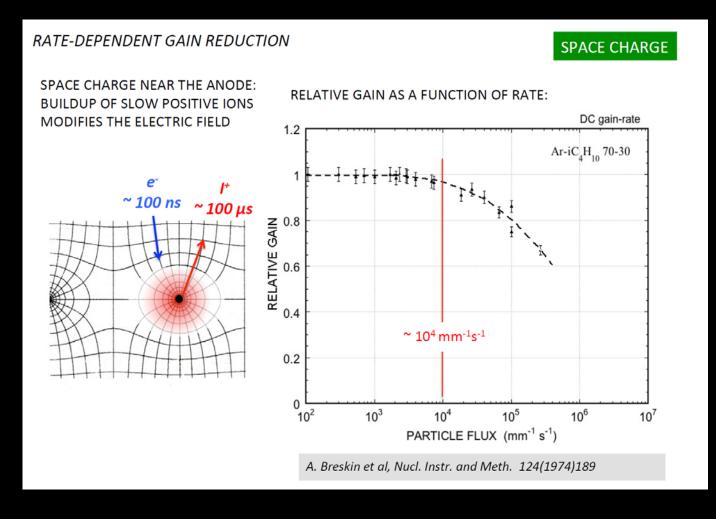


Review on TPC's

- TPC is a 3-D imaging chamber
 - Large dimensions. Little material
 - Slow device (~50 μs)
 - 3-D coordinate measurement ($\sigma_{xy} \approx 170~\mu\text{m}$, $\sigma_z \approx 740~\mu\text{m}$)
 - Momentum measurement if inside a magnetic field
- Reviewed some the main ingredients
 - Field cage, gas, wire chambers, gating grid, laser calibration, electronics, etc.
- History
 - First proposed in 1976 (PEP4-TPC)
 - Used in many experiments
 - Well established detecting technique

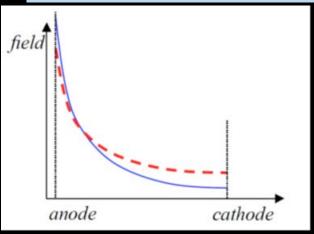


Limitation factors of MWPC: effects on the rate capability



Despite their successful use in particle physics experiments and other fields, MWPCs have several limitations, intrinsic in their conception. The creation in the multiplication process of large amounts of positive ions, slowly receding towards the cathodes, causes a modification of the applied electric field, and results in a drop of gain and efficiency at particle fluxes above ~10⁴mm⁻¹s⁻¹.

The discrete wire spacing is itself a limitation to the multi-track resolution, essential at high particle rates and multiplicities.

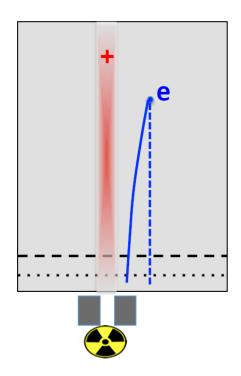


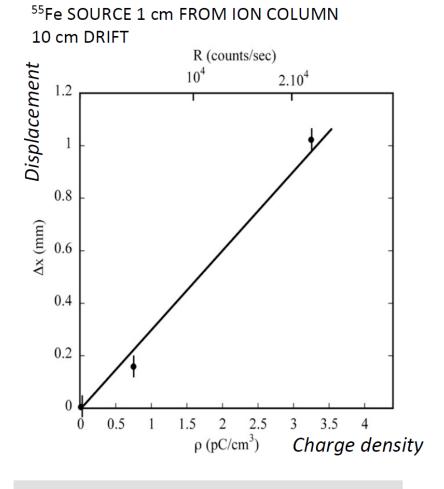
Limitation factors of MWPC: effects on the spatial coordinates determination

POSITIVE ION BACKFLOW

SPACE CHARGE

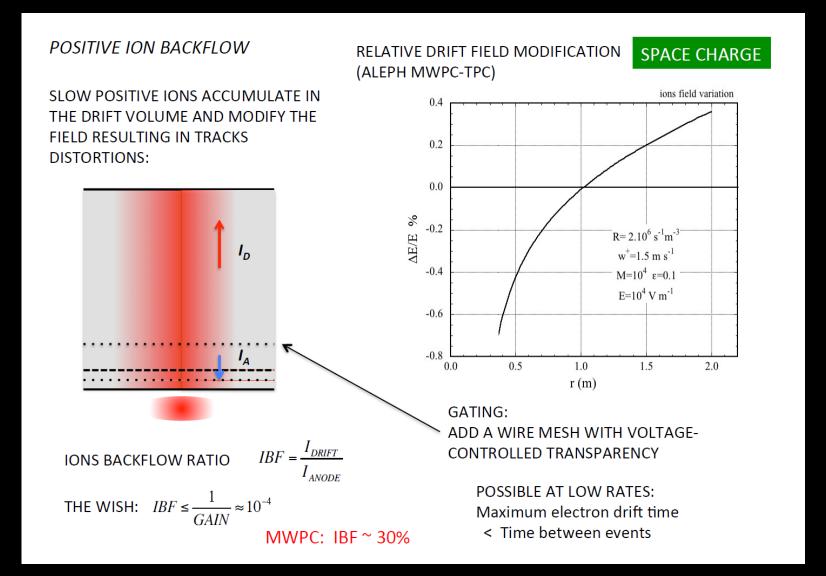
LATERAL DISPLACEMENT OF ELECTRONS
DRIFTING NEAR A POSITIVE IONS COLUMN

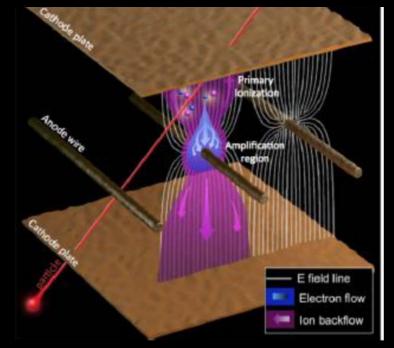




D. Friedrich, et al, Nucl. Instr. and Meth. 158(1979) 81

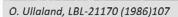
Limitation factors of MWPC:

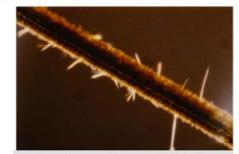




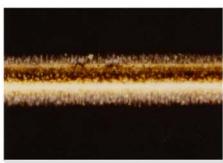
Limitation of Gas Detectors: the ageing

AGING SECONDARY PROCESSES Polymerization of organic compounds with formation of deposits on thin wires:





I. Juric and J. Kadyk, LBL-21170 (1986)141



I. Juric and J. Kadyk, LBL-21170 (1986)141



M. Binkley et al, Nucl. Instr. and Meth. A515(2003)53

Even more detrimental, the creation and deposit on the anode wires of thin insulating layers caused by the polymerization of organic gases or pollutants may result in an amazingly short operating life span

Classical ageing

Avalanche region → plasma formation (complicated plasma chemistry)

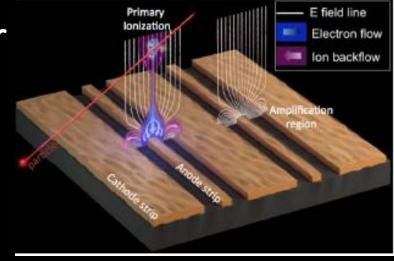
- Dissociation of detector gas and pollutants
- Highly active radicals formation
- Polymerization (organic quenchers)
- Insulating deposits on anodes and cathodes



Anode: increase of the wire diameter, reduced and variable field, variable gain and energy resolution.

Cathode: formation of strong dipoles, field emmision and microdischarges (Malter effect).

An innovative structure of Gas Detector



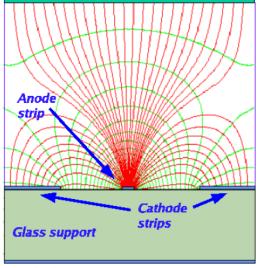
MICRO-STRIP GAS COUNTER

MICRO-PATTERN GAS DETECTORS

MSGC

Anton Oed, 1988

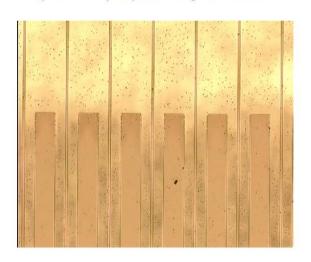
Drift electrode



Back plane

A. Oed, Nucl. Instr. and Meth. A263(1988)351

10 μm wide anode strips, 50 μm cathode strips at 100 μm pitch on glass substrate:

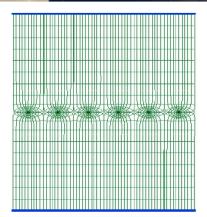


The Micro-Strip Gas Counter (MSGC), introduced by Anton Oed in 1988, seemed to overcome some of the above mentioned limitations. Consisting of a set of thin parallel metallic strips laid on an insulating substrate, alternatively connected as anodes and cathodes, MSGCs provide rate capabilities two orders of magnitude higher than MWPCs, and a tenfold improvement in the multi-track resolution. Disappointingly and despite the efforts by many groups, the device appeared to be rather susceptible to irreversible degradation due to occasional but destructive discharges.

To compare a bit MWPC and MSGC

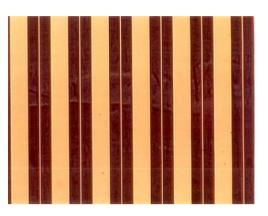
MWPC

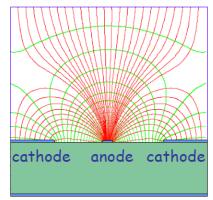
CHA TABLER



Typical distance between wires limited to 1 mm due to mechanical and electrostatic forces

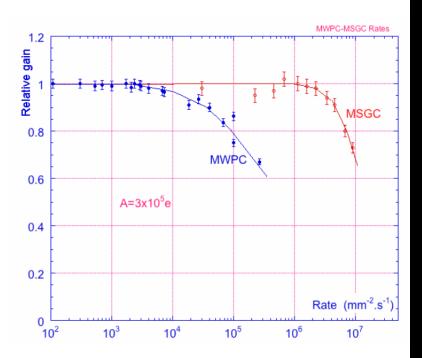
MSGC





Typical distance between anodes 200 μm thanks to semiconductor etching technology

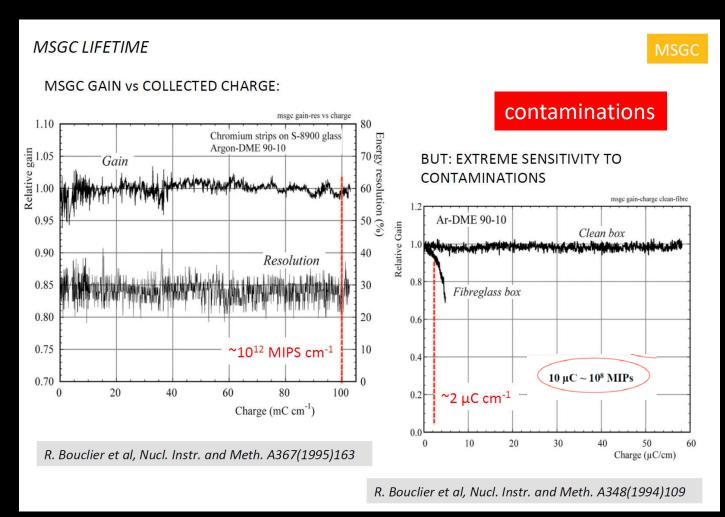
A. Oed Nucl. Instr. and Meth. A263 (1988) 351.

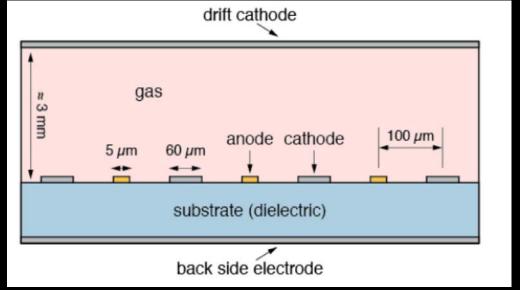


Rate capability limit due to space charge overcome by increased amplifying cell granularity

R. Bouclier et al, Nucl. Instr. and Meth. 367(1996)328

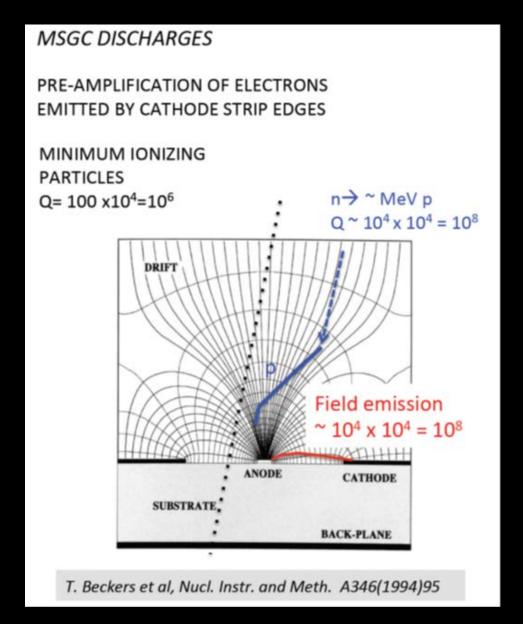
A few problems with MSGC: contamination



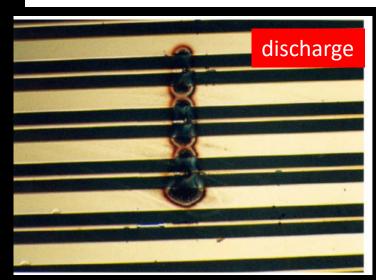


The problems met with the MSGCs resulted in a large effort devoted to the development of sturdier structures, preserving its rate and multitrack capabilities

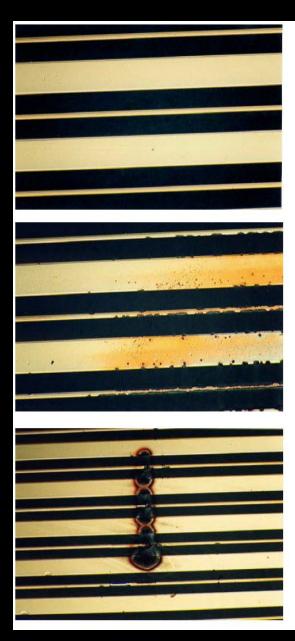
A few problems with MSGC: discharge issue



Many of the problems encountered with MSGCs are connected to the use of fragile electrodes exposed to the high electric fields needed to achieve the gains, typically around 10⁴, needed for detection of small ionization yields. Under these condition,the occurrence in the gas of rare but highly ionizing events, due for example to neutron or gamma conversions, may lead to the creation of a local charge density exceeding the Raether limit (10⁷ electron—ion pairs) leading to the formation of a streamer, and eventually to a discharge.



Discharge vs gain in MSGC



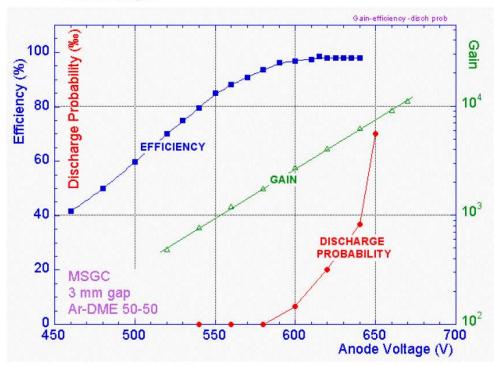
Surface charging

Bulk resistivity of the support material Surface modification by doping or deposition

Ageing

Gas, Gas system, MSGC support, Construction material

Discharges



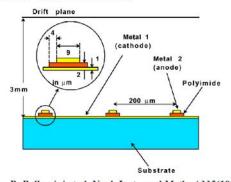
New MPGDs (exploiting micro-litographic technique)

Semiconductor industry technology:

Photolithography Etching Coating

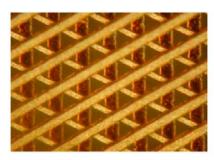
Doping

MICRO-GAP CHAMBER



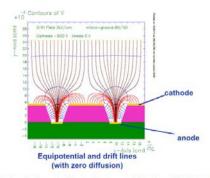
R. Bellazzini et al, Nucl. Instr. and Meth. A335(1993)69

FIELD GRADIENT LATTICE DETECTOR (FGLD)



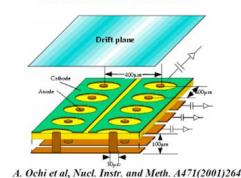
L.Dick et al Nucl. Instr. and Meth. A535(2004)347

MICRO-GROOVE CHAMBER



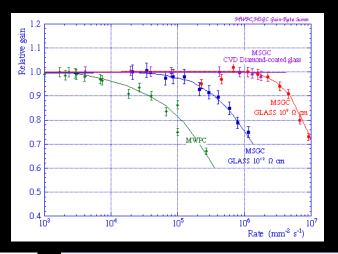
R. Bellazzini et al, Nucl. Instr. and Meth. A424(1999)444

MICRO-PIXEL CHAMBER



Advantages of MPGDs

- 1. Rate Capability
- 2. High Gain
- 3. Space Resolution
- 4. Time Resolution
- 5. Energy Resolution
- 6. Ageing Properties
- 7. Low Material Budget
- 8. Geometrical Flexibility
- Readout Structures
- 10. Ion Backflow Reduction
- 11. Photon feedback Reduction
- 2. Large area at low prices
- 13. Flexible geometry

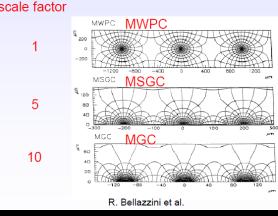


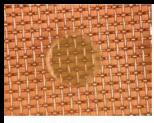
Problem:

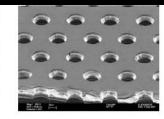
 rate capability limited by space charge defined by the time of evacuation of positive ions

Solution:

 reduction of the size of the detecting cell (limitation of the length of the ion path) using chemical etching techniques developed for microelectronics and keeping at same time similar field shape.







GEM



Micromegas

THGEM

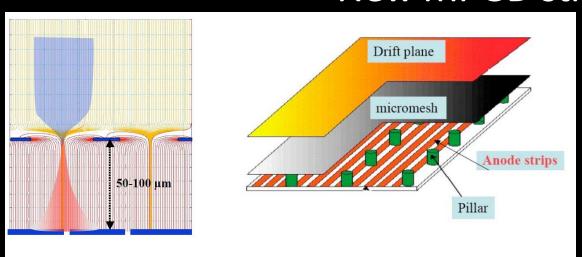
Current Trends in Micropattern Gaseous Detectors

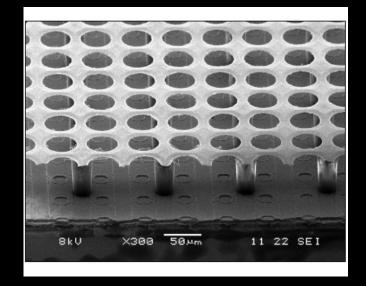
- 1. Manufacturing Technologies
- 2. Micromegas
- 3. GEM
- 4. ThickGEM/RETGEM
- 5. MPDG with CMOS pixel ASICs
- 6. Ingrid Technology

Applications (HEP, Astrophysics, Nuclear Physics, Industrial and Medical)

- 1. Charged Particles Tracking
- 2. Triggering
- 3. TPC Readout
- 4. Calorimetry, Muon Detectors
- 5. Photon Detectors (UV and Visible Light Detection)
- 6. X-Ray Astronomy
- 7. Soft X-Ray Imaging
- 8. Neutron Detection
- 9. Cryogenic Detectors

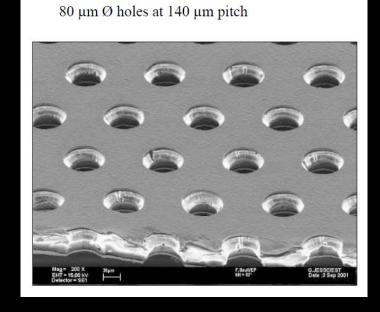
New MPGD structures

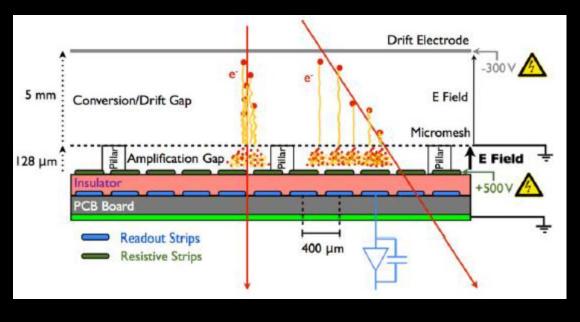




Y. Giomataris et al, Nucl. Instr. and Meth. A376(1996)29

Holes pattern on Cu-plated polymer 50 μm thich 10-150 μm



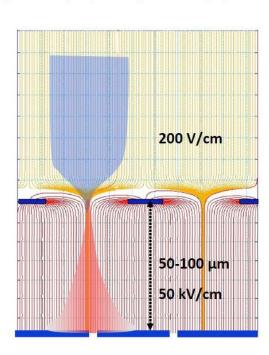


The structure of MMEGAS

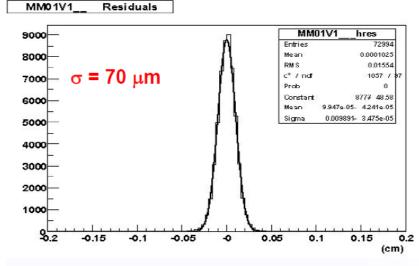
NEW MICRO-PATTERN GAS DETECTORS

MICROMEGAS

MICROMEGAS
Thin (50-100 μm) multiplication gap:



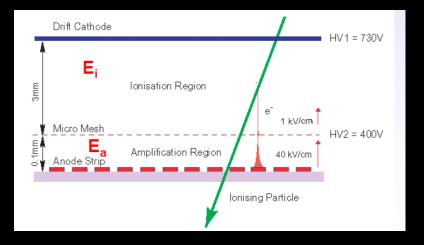
Y. Giomataris et al, Nucl. Instr. and Meth. A 376(1996)29

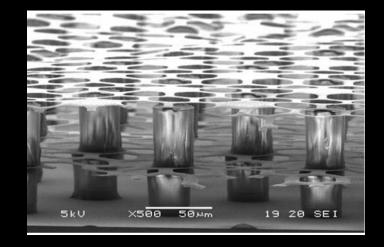


Space resolution

E field similar to parallel plate detector. $E_a/E_i \sim 50 \text{ to secure electron transparency} \\$ and positive ion flowback supression.

J. Derré et al, Nucl. Instr. and Meth. A459(2001)523



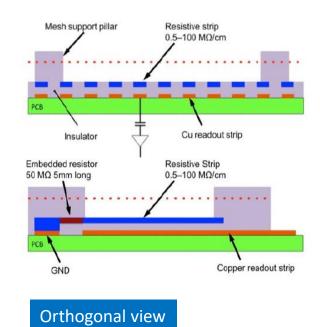


Discharge mitigation technique in MMEGAS

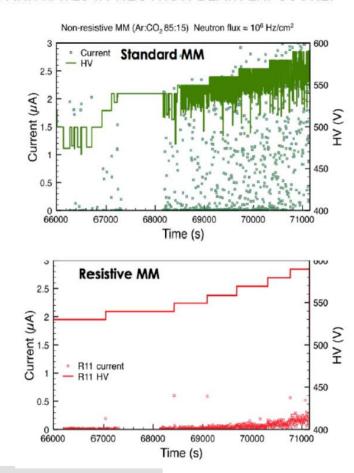
DISCHARGES: RESISTIVE MICROMEGAS

MICROMEGAS

BUILT ON A HIGH-RESISTIVITY POLYMER



SPARK RATES IN NEUTRON BEAM EXPOSURE:



T. Alexopolous et al, Nucl. Instr. and Meth. A640(2011)110

Gas Electron Multiplier detectors

From 1997 MPGDs have played a fundamental role in **HEP** and Nuclear Physics. Today their applications are being extended beyond fundamental Science:

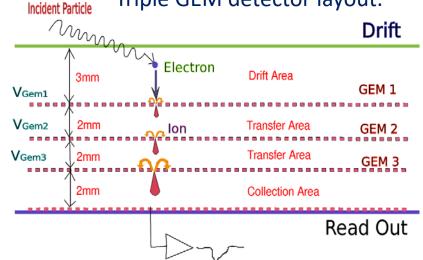
Astrophysics
Material analysis
Heritage and Art
Homeland security
Industrial
Risk alert

Medicine

Environment

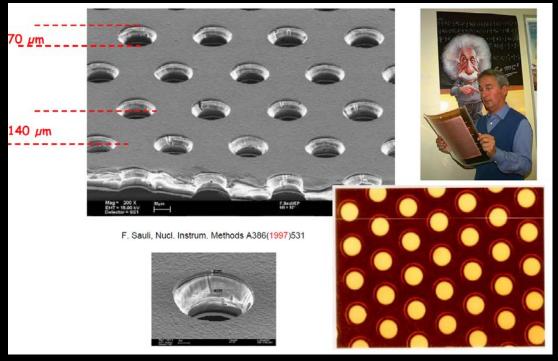
77.827-818 817.827-917-91 GEM foil.

Triple GEM detector layout.

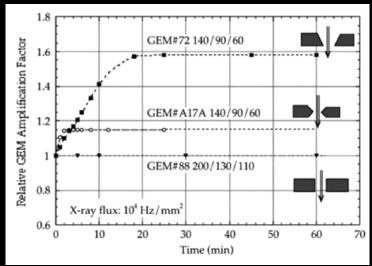


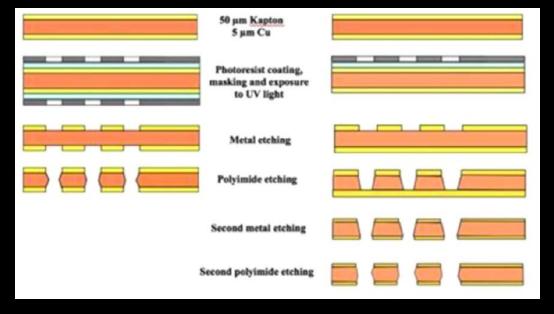
- √ high detection efficiency (>98%);
- ✓ good 2D imaging capability by read-out segmentation;
- ✓ excellent spatial resolution typically 100 μm;
- √ good time resolution (4-5 ns);
- √ high radiation flux resistance;
- √ high rate capability (50 MHz/cm²)
- ✓ flexibility, robustness and low costs.

GEM foil construction technology



Thin-metal coated polymer foil Pierced by a high density of holes (50-100/mm²). Typical geometry: 5μm Cu on 50μm Kapton, 70μm holes with 140μm pitch





Double mask and single-mask tecnology

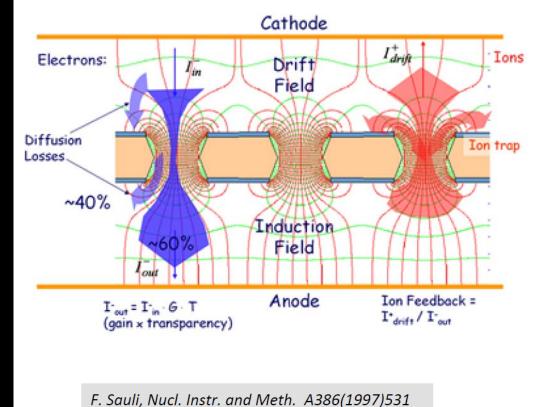
Single GEM structure

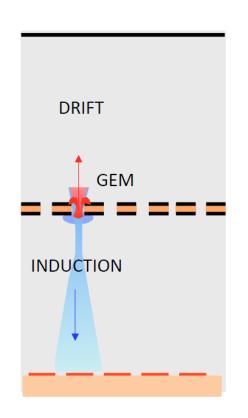
GAS ELECTRON MULTIPLIER (GEM)

GEM

Thin (50 μ m) metal-coated polymer foil with high density of holes:

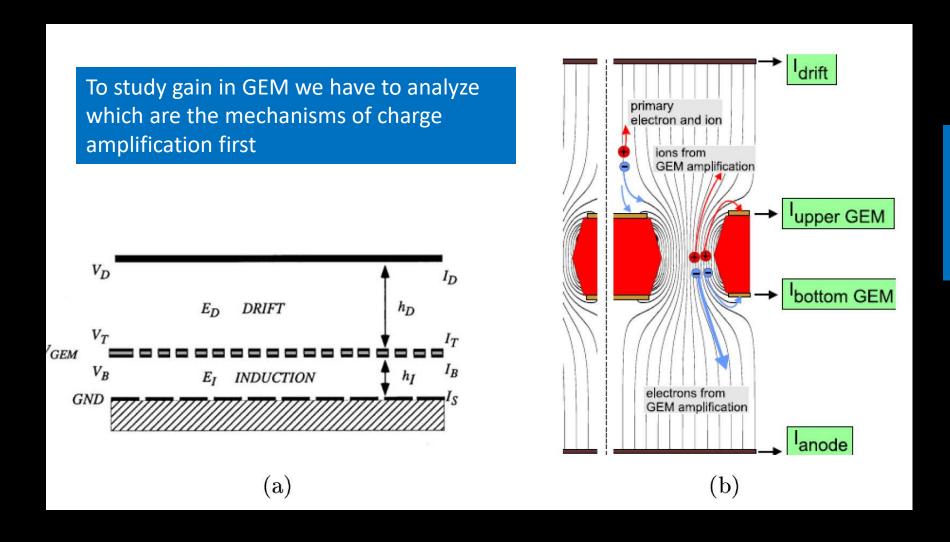
FAST ELECTRON SIGNAL ON ANODE STRIPS (NO ION TAIL):





- 1. Electrons are collected on patterned readout board.
- 2. A fast signal can be dected on the lower GEM electrode for triggering or energy discrimination.
- 3. All readout electrodes are at ground potential.
- 4. Positive ions partially collected on the GEM electrode

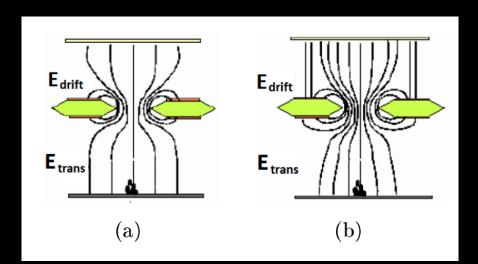
The choice of parameters in a single GEM foil detector

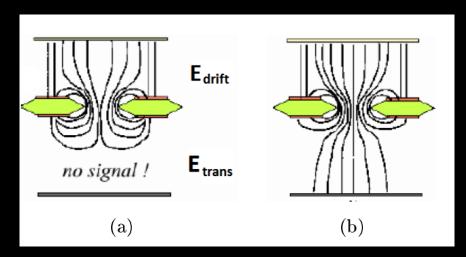


We identify three electric fields:

The drift field, the moltiplication field and the induction field

The electron Transparency





- 1. The electric fields (drift and induction) play a crucial role to define the so called *Transparency*
- 2. Another important parameter is the geometry of holes



Focusing (a) or undefocusing (b) effect on electron determines the *sticking* ε_{coll} and the *transfering* ε_{extr} efficiencies

The intrinsic gain of GEM foil can be defined as:

$$G_{intr} \propto e^{\langle \alpha \rangle V_{GEM}}$$

The effective gain is defined as:

$$G_{eff} = G_{intr} \bullet \varepsilon_{coll} \bullet \varepsilon_{extr}$$

Rigorous formula of Gain

$$G = \exp(\int (\alpha(x) - \eta(x))\delta x),$$

Gain determination

$$\epsilon^{coll} = \frac{electrons\ collected\ in\ the\ holes}{electrons\ produced\ above\ the\ holes}$$

Collection efficiency

$$\epsilon^{extr} = \frac{electrons\ extracted\ from\ the\ holes}{electrons\ produced\ in\ the\ holes}$$

Extraction efficiency

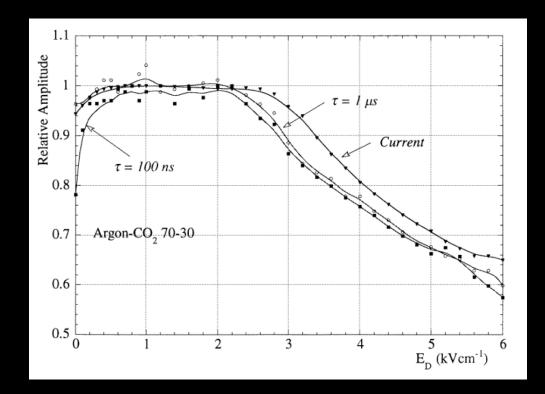
$$G_{int} \propto e^{\overline{\alpha} \sum V_{GEM}}$$

Intrinsic gain

$$G_{eff} = G_{intr} * \varepsilon^{coll} * \varepsilon^{extr} = G_{intr} * T$$

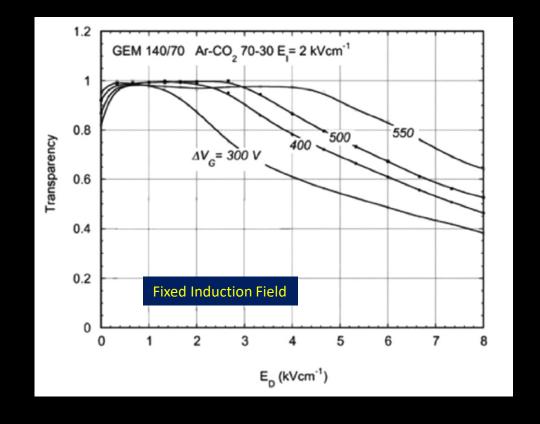
Effective gain

Influence of Electric Fields on the electron transparency

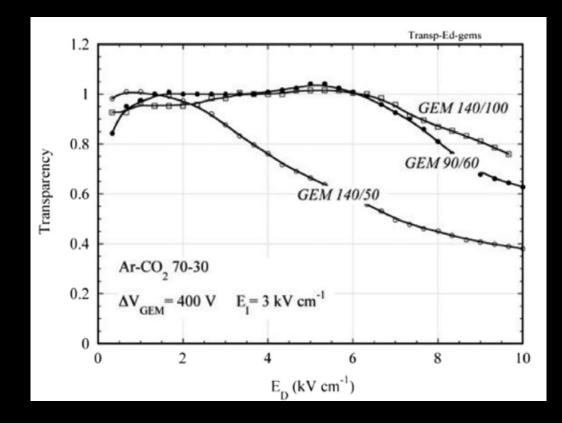


Owing to the structure of the detector, the sharing of collected charges (electrons and ions) between electrodes depends on the value of fields and holes geometry characteristics

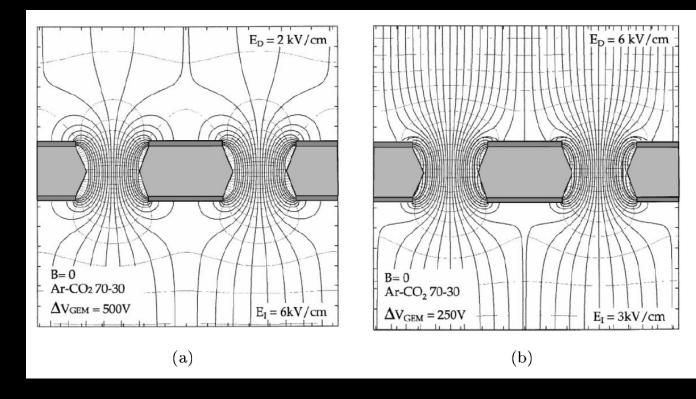
Electron transparency of a standard GEM electrode (70 um holes at 140 um pitch) as a function of drift field for fixed induction field, for several values of GEM voltage.



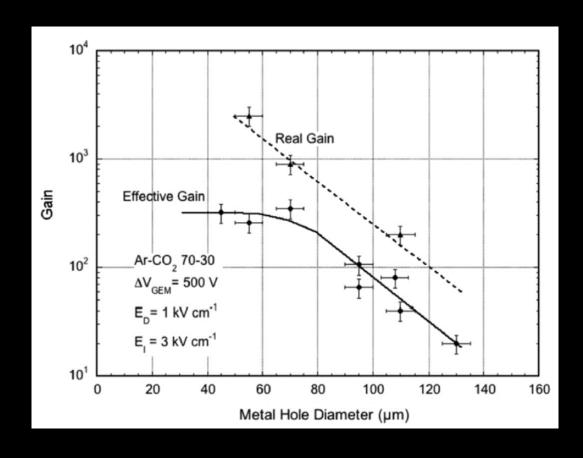
Influence of Electric Fields on the electron transparency



Electron transparency dependence vs Different drift and induction electric field Electron transparency for different GEM geometry labeled as pitch/holes diameter.



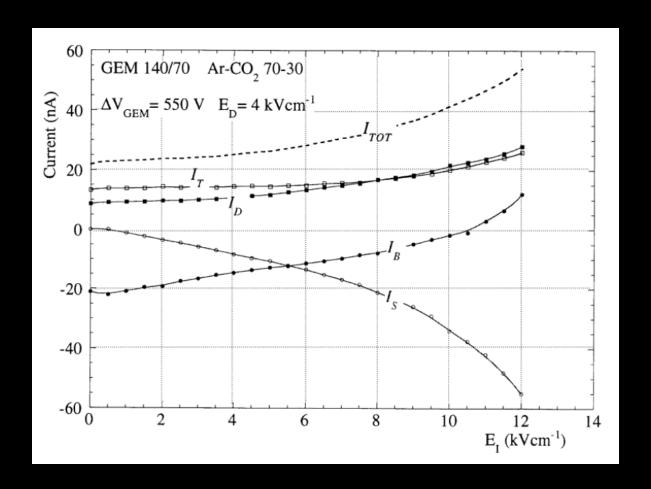
Effect of hole dimension on the Gain

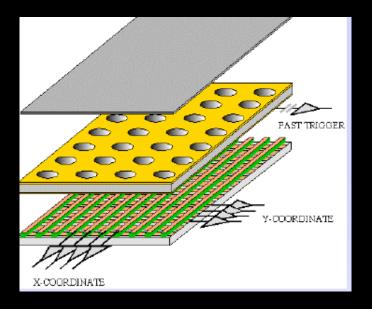


Reduce too much the hole diametr is not beneficial on the gain value

Choice of drift and induction gaps thicknesses: current signal on a single GEM detector

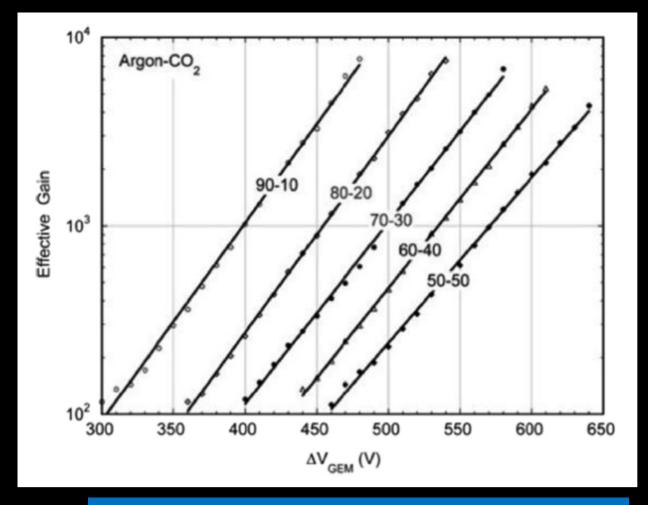
The drift region thickness is chosen such to guarantee an appropriate number of primary ionization pairs and to reduce pile-up event in order to have a good event rate





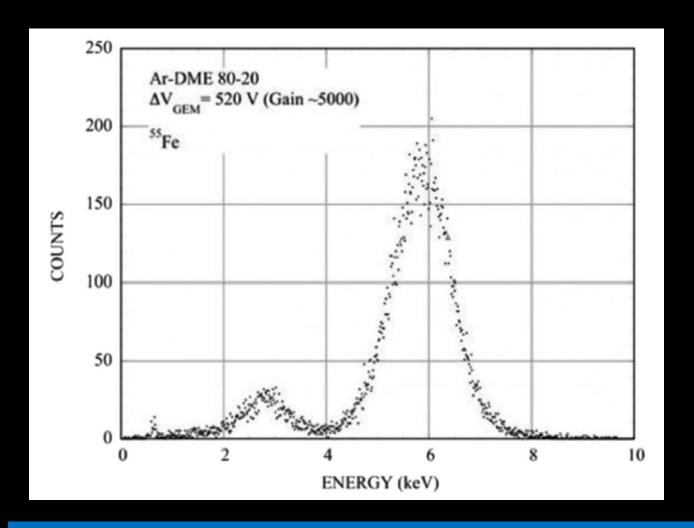
The induction gap thickness is such that to avoid the formation of sparks

Gain dependence on gas mixture composition



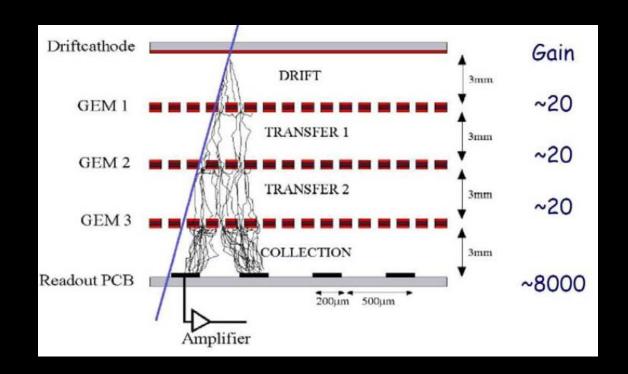
Single GEM effective gain as a function of voltage in Ar–CO₂ mixtures at atmospheric pressure as a function of HV

Energy resolution for a single GEM detectors



Pulse height spectrum on 5.9keV for a single GEM. The relative energy resolution is 17%FWHM.

Effective Gain in Triple-GEM detector



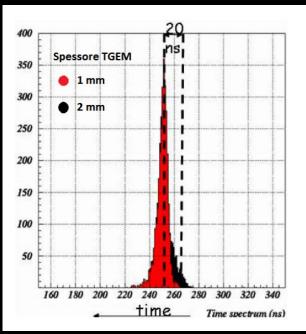
- Multiple structures provides equal gain at lower voltage
- Discharge probability on exposures to α particles is strongly reduced

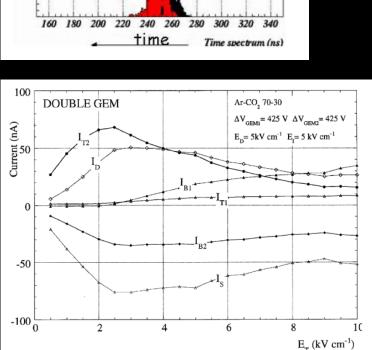
$$G_{intr} \propto e^{<\alpha>V_{GEM_{TOT}}}$$

$$G_{eff} = \prod_{i=1}^{3} G_{intr}^{i} \cdot \epsilon_{infil}^{i} \cdot \epsilon_{estr}^{i}$$

$$\epsilon_{infil}^{i}\!\cdot\!\epsilon_{estr}^{i}=T^{i}$$

Dimensions of different regions









Important on the first transfer gap thickness

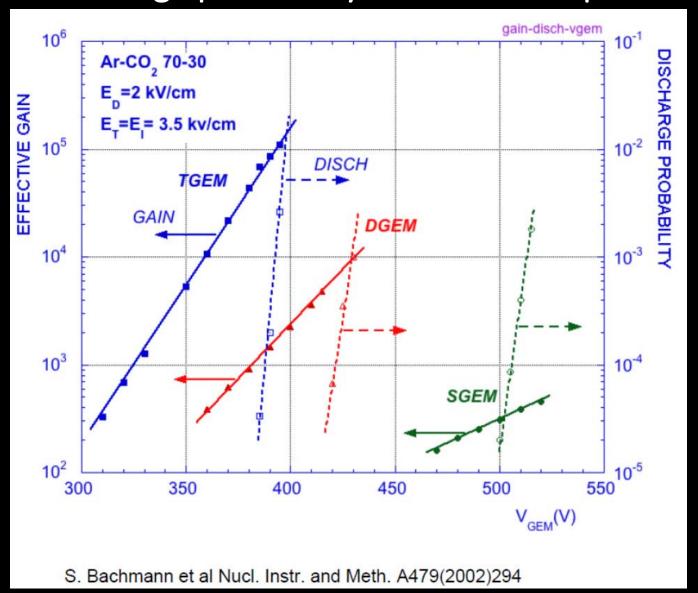
Limitations of sparks



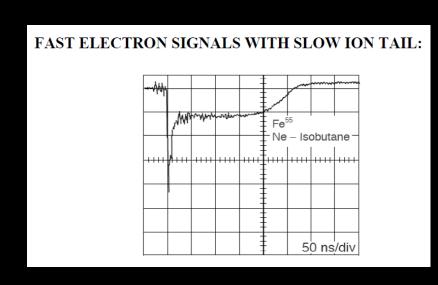
Important on the second transfer gap thickness

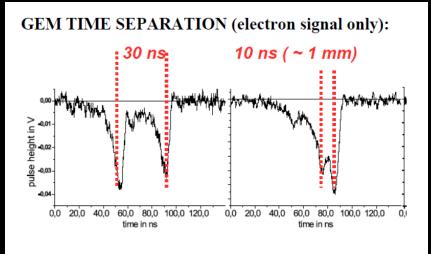
Dependence of current signal on Transfer Electric Field in a double-GEM

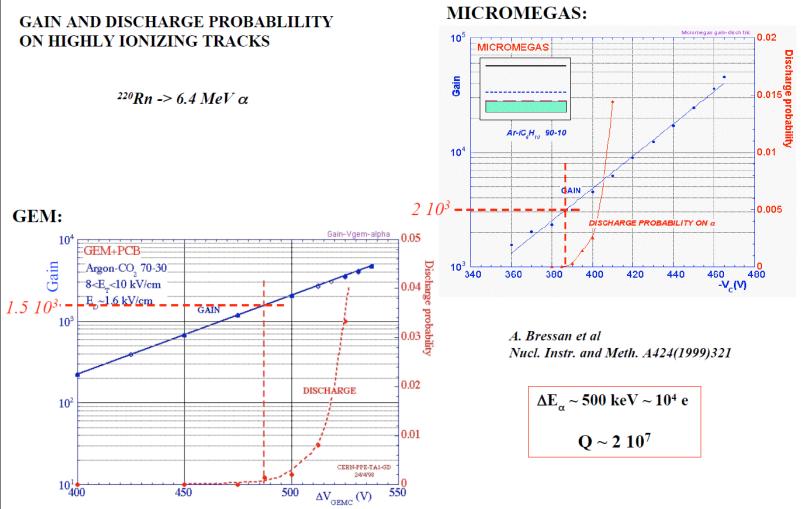
Multi-GEM detector Gain and discharge probability with 5 MeV α particles



Performances comparison MMEGAS vs GEM



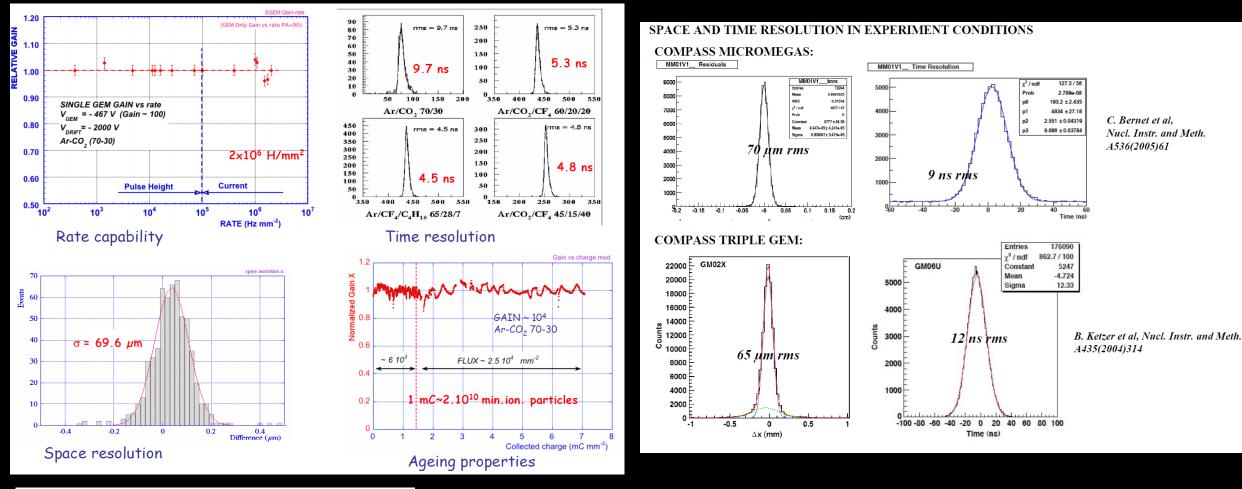


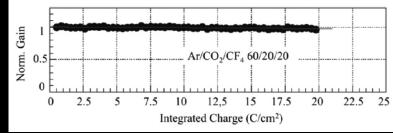


Discharge probability vs effective gain

DISCHARGES: MULTI-GEM MULTI-GEM DISCHARGE PROBABILITY VS GAIN: IN MULTI-GEMs, THE CHARGE SPREADS BY **DIFFUSION OVER MANY INDEPENDENT HOLES!** Disch S-D-TGEM 5 10 Discharge probability 2 10-3 Q7108 Q~109 Triple GEM Q~107 **Double GEM** Single GEM 1 10-3 0 10² 10³ 10⁵ 10⁴ 10⁶ **Effective Gain**

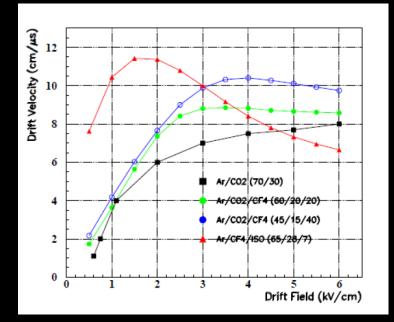
Standard GEM typical spatial & timing resolution

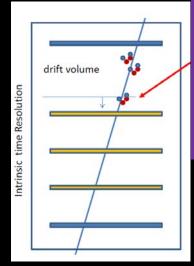


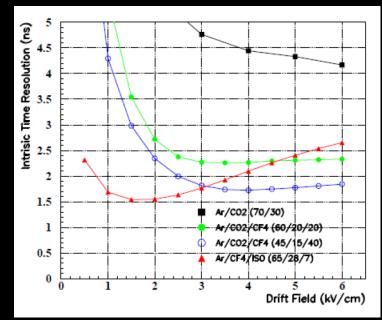


Extended lifetime with addition of CF₄ to standard mixture Ar/CO2

Timing performances







the time resolution is dominated by fluctuations of the nearest distance of the primary ionisation processes to the region where the gain is acquired, dnear.

Defining λ as the average number of primary clusters generated by an ionising particle inside the gas, this distance follows a classical exponential distribution

$$d = exp(-\lambda x)/\lambda$$

The **contribution** to the time resolution of the **drift velocity** is:

$$\sigma_t = (\lambda v_d)^{-1}$$

With a drift gap of the order of 3-4 mm and with a proper choice of the gas mixture, these detectors can reach a time resolution of the order of 5-10 ns.

Typical values for gases employed in MPGDs are $\lambda = 3 \text{ mm}^{-1}$ and v_d up to 0.12 mm/ns leading to few **ns time resolution** with the best choice of gas mixtures and operating voltages

Timing performances

...moreover the contribution of the gain fluctuation is governed by the gas time constant (effectiveTownsend coefficient)

(NeffVd)-1

In the multiplication volume the field is very high (50 -100 kV/cm) and consequently $\eta_{\rm eff} > 100 \ {\rm mm}^{-1} \ {\rm e} \ {\rm con}$ $v_d = 0.12 \ {\rm mm/ns} \ \rightarrow {\rm time} \ {\rm resolution} \ {\rm contribution} \ < 100 \ {\rm ps}$

In order to improve the time resolution new configurations are proposed. The improvement is obtained by using several drift regions each one coupled to its multiplication stage

Preliminary Events/2ns Drift electrode Entries χ^2 / ndf 2.4 / 4 Prob $5.9e+02 \pm 6.2e+01$ Mean 25 ± 0.2 Sigma 1.7 ± 0.1 Pion Beam Both Lavers powered Drift Fields = 8 kV/cm Amplification Fields = 120 kV/cm Gas Mixture = Ar/CO₂ 70/30 Signal pickup from drift electrode Time(ns)

I. Vai et.al. CMS Collaboration Vienna Conference on Instrumentation2016

The reduction of the time resolution, in fact, is proportional to the number of the layers *N*_D employed. In this case the time resolution transforms in:



$$\sigma_t = (\lambda v_d N_D)^{-1}$$

R. De Oliveira et.al. «A novel fast timing micropattern gaseous detector:FTM» CERN-OPEN 2015-002 INFN-15-01/BA

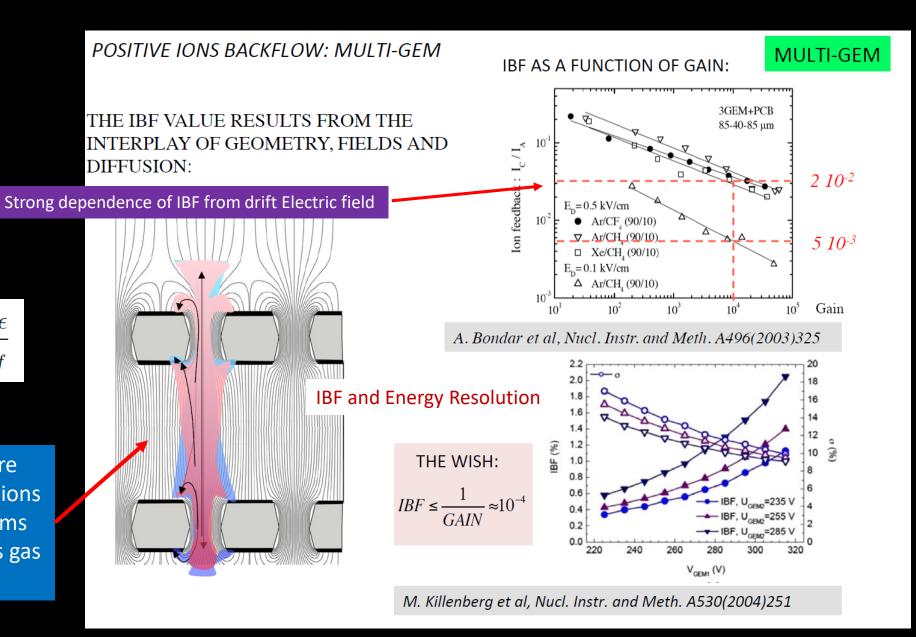
IBF effect in multi-GEM

$$IBF = \frac{N_C^+}{N_A^-}$$

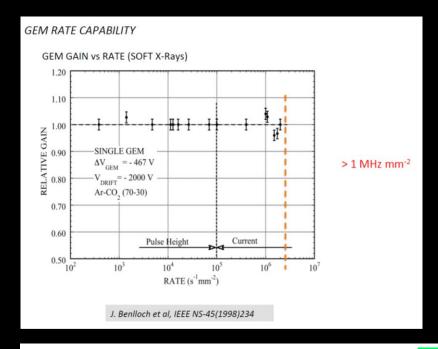
$$\epsilon = \frac{N_G^+}{N_I}$$

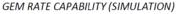
$$IBF = \frac{N_I + \epsilon N_I}{N_I G_{eff}} = \frac{1 + \epsilon}{G_{eff}}$$

The extraction of electrons is more efficient of the recombination of ions in the drift region. This mechanisms is favoured in GEM than in others gas detectors

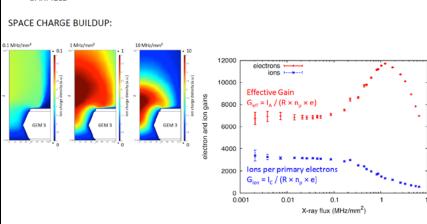


Space Charge effect on GEM rate capability



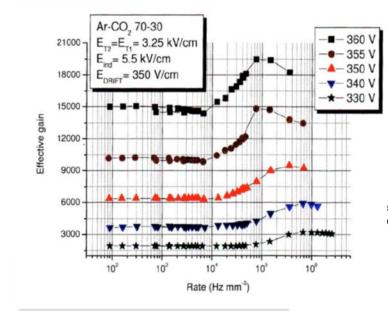


- COMSOL Finite Element Analysis
- GARFIELD





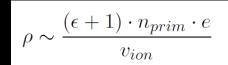
A STRANGE OBSERVATION: GAIN INCREASE AT VERY HIGH RATES (2006, UNPUBLISHED)



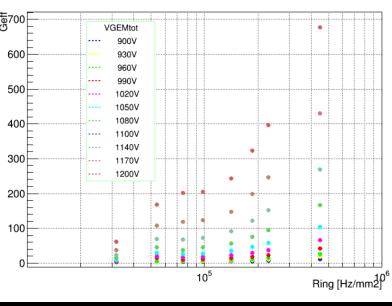
Peter Everaerts, PhD Thesis Gent University (2006)

$$E(z) \sim E_D - \frac{\rho d}{2\epsilon_0} + \frac{\rho z}{\epsilon_0}$$

Electric field in drift region on the top of 1'st GEM

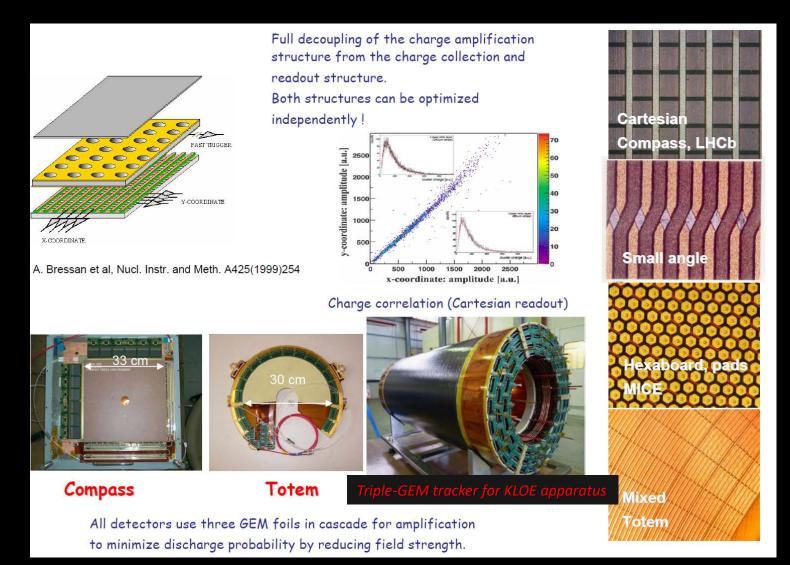


Spatial charge density



The IBF study on a TPG prototype, Thesis_2016 Bari University

GEM different shapes and readout board





- Development of GEM detector of large area
- Triple GEM for CMS-End Cap Muon detector

Gas Detestors in LHC experiments

ALICE: TPC (tracker), TRD (transition rad.), TOF (MRPC), HMPID (RICH-pad chamber),

Muon tracking (pad chamber), Muon trigger (RPC)

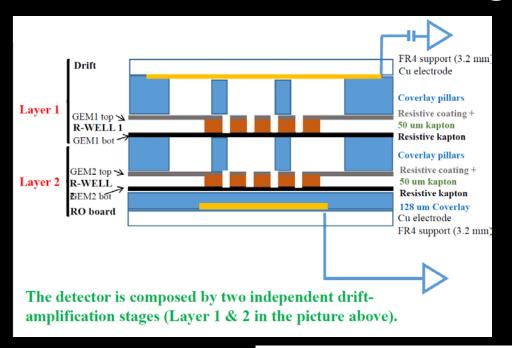
ATLAS: TRD (straw tubes), MDT (muon drift tubes), Muon trigger (RPC, thin gap chambers)

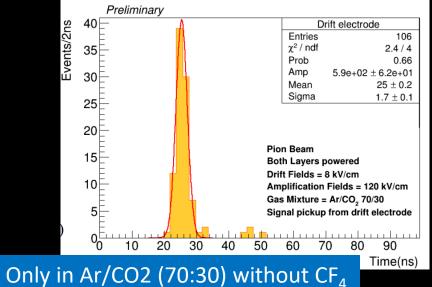
CMS: Muon detector (drift tubes, CSC), RPC (muon trigger)

LHCb: Tracker (straw tubes), Muon detector (MWPC, GEM)

TOTEM: Tracker & trigger (CSC, GEM)

Fast Timing architecture





Each amplification region is based on *a pair of polyimide foils* stacked due to the electrostatic force induced by the polarization of the foils:

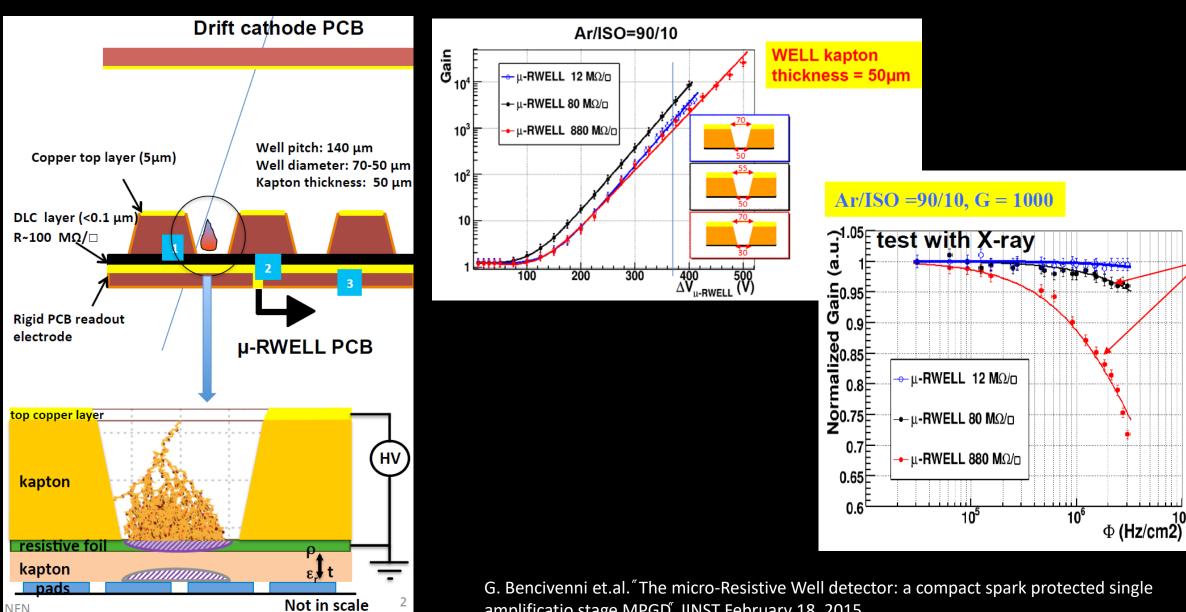
- The first foil, perforated with inverted truncated-cone-shaped holes (bases 100 μm and 70 μm, pitch 140 μm), is a 50 μm thick Apical KANECA, coated with diamond-like carbon technique, to reach up to 800 MΩ/□ resistivity.
- The second foil is 25 μm thick XC Dupont Kapton, with a resistivity of 2 $M\Omega/\Box$.

The *drift volumes are 250 \mu m thick*, with planarity ensured by coverlay pillars, with diameter 400 μm and pitch of ~3.3 mm.

The active area (circular) is about 20 cm².

D. Abbaneo et.al., "R&D on a new type of micropattern gaseous detector: The Fast Timing Micropattern detector", arXiv:1503.05330v1 European Patent Application 14200153.6 M. Maggi, A. Sharma, R. De Oliveira

Novel MPGD architecture

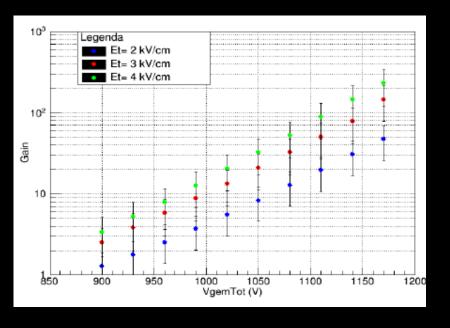


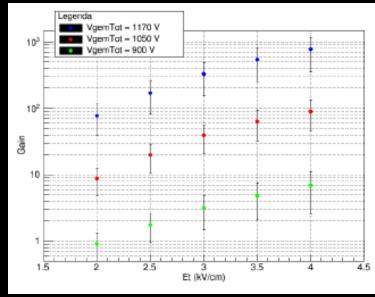
amplificatio stage MPGD, JINST February 18, 2015

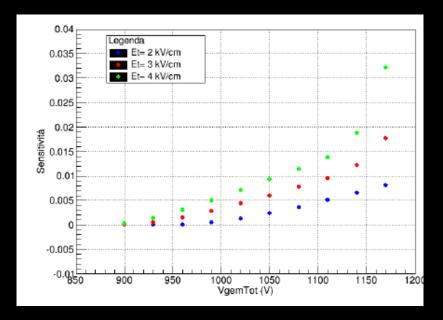
Laboratory measurements

- 1. Gain vs $V_{GEM TOT}$ for two gas mixtures
- 2. Transparency measurement:
 - @fixed V_{GFM} Gain vs E_d for different gas mixtures
 - Sensitivity measurements

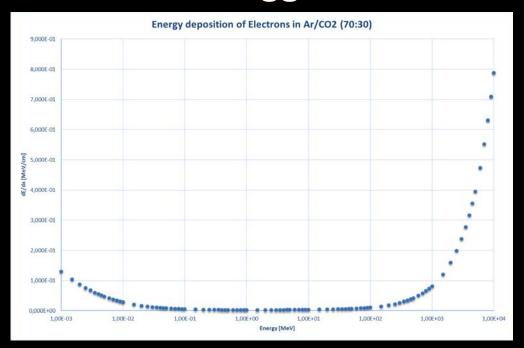
$$G = \frac{I_a}{R \times n_T \times e}$$

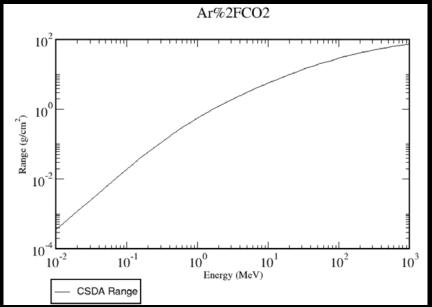






Suggestion for Gain measurements





Evaluation of primary charge

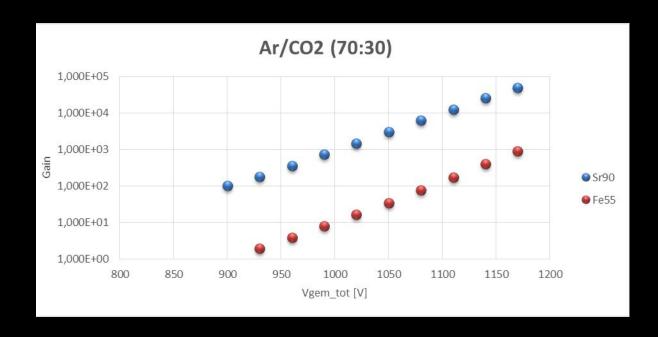
Fe⁵⁵ X-ray of 5.9 KeV → photoelectric effect

- → photoelectrons completely converted in drift gap
- → Range < 0.3 cm of drift gap

$$n_T = \Delta E \times \left[\frac{0.7}{W_i(Ar)} + \frac{0.3}{W_i(CO_2)} \right] = 5900 \times \left[\frac{0.7}{26} + \frac{0.3}{33} \right] \approx 212$$

Sr⁹⁰ β of 0.546 MeV and 2.28MeV \rightarrow range » drft gap \rightarrow we may assume $\langle N_T \rangle = 28 e^-$

Space charge effect



Triple-GEM detector 10x10 cm²

Configuration: 3-1-2-1

Operated with:

• Ed = 1.5 KV/cm

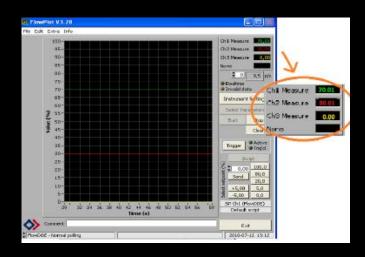
• Et1 = Et2 = 3.0 KV/cm

• Ei = 5 KV/cm

 \bullet $V_{G1} > V_{G2} > V_{G3}$

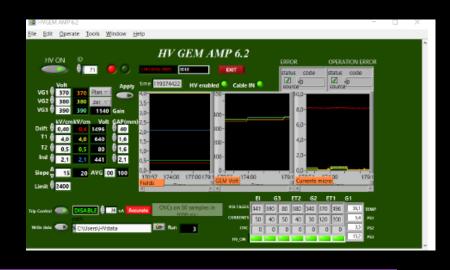
Experimental setup





Triple gas system controlled by Bronkhorst Flowmter

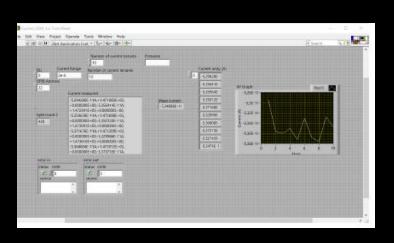




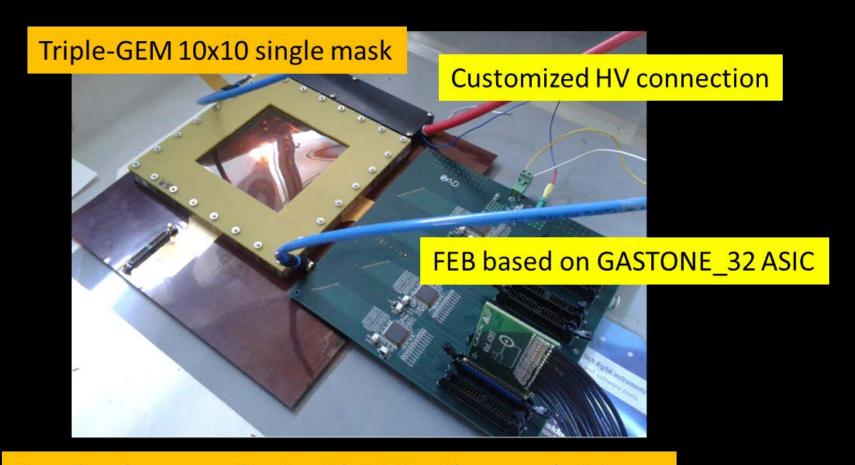
Special purpose custom-made HV system



Pico-ammeter Keithley 6487



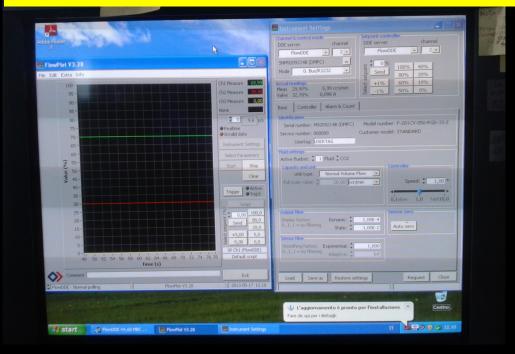
The triple-GEM detector



Anode plane readout with X-Y orthogonal strips $600\mu m$ width, $100\mu m$ pitch

GEM-Lab set-up

(FLOWDDE/FLOWPOT Flow-meter control software)



(EL-FLOW® Bronkhorst High Tech BV)



- •The gas system is composed by three mixed lines (Ar-CO2-CF4)
- •Each line is controlled by three Flow-mass meters

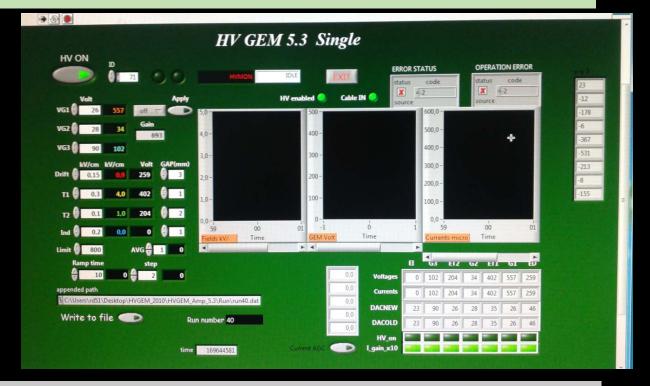
Ar (Max Capability = 20 cc/min), CO2 (20 cc/min), CF4 (2 cc/min) (precision of 1%)

•All the system is controlled by a PC software through an RS232 serial line interface

GEM-Lab set-up



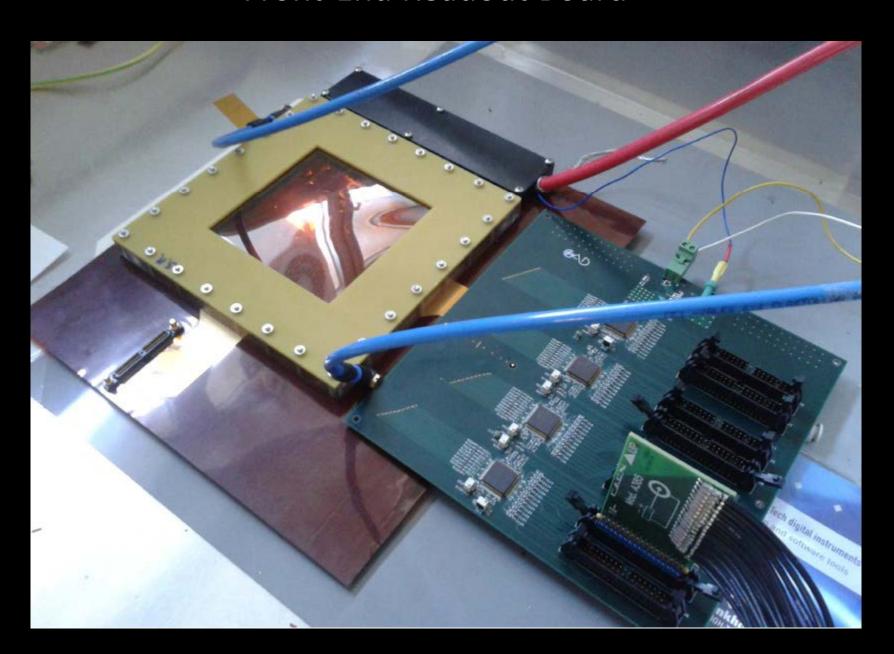
- •HV power supply based on Frascati HVGEM module and used in LHCb experiment (last release with Trip control on each channel
- •6 channels 150 μA-750 V_{MAX} /channel
- •1 channel 75μA 1500 V_{MAX}
- •HVGEM control is made through a LabView program
- •Interfaced to PC through a USB connection



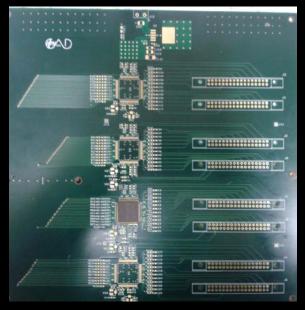
G. Corradi, F. Murtas and D.Tagnani, A novel High Voltage System for a triple GEM detector.

Nuclear Inst. and Methods in Physics Research, A (NIMA46 128)

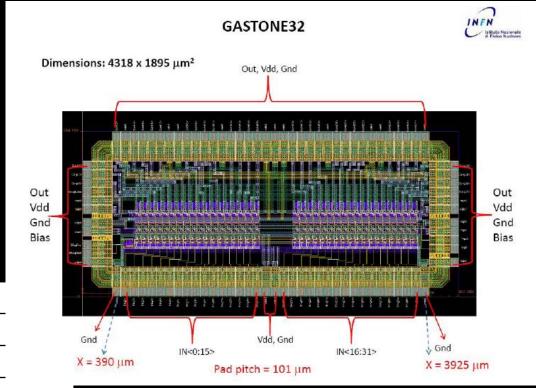
Front-End Readout Board



GASTONE_32 main features



Input impedance	120 Ω
C _{det} range	0 – 150 pF
Number of channels	32
Baseline restorer	No
Total Gain sensitivity	20 mV/fC ($C_{det} = 0 pF$)
Peaking time	90 ns (C _D =100 pF)
Measured X-talk	< 1%
ENC (rms) measured on detector	800 e ⁻ + 40 e ⁻ /pF
Input protection circuitry	Integrated in each input channel
Power consumption	5 mW/Ch

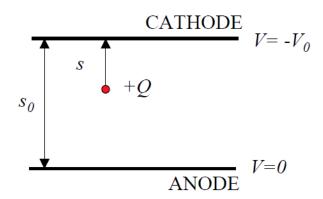


- •Capable to drive a 50Ω line
- •Signal referred to GND

Backup

Charge induction – ionization chamber

SIGNAL DEVELOPMENT BY A MOVING CHARGE +Q



Electrons- ion pair (-Q and +Q) released at the same distance s from the cathode :

$$q(t) = Q\left(\frac{w^{-}t}{s_0} + \frac{w^{+}t}{s_0}\right) \quad 0 \le t \le T^{-}$$

$$q(t) = Q\left(\frac{s - s_0}{s_0} + \frac{w^{+}t}{s_0}\right) \quad T^{-} \le t \le T^{+}$$

$$q(T^{+}) = Q$$

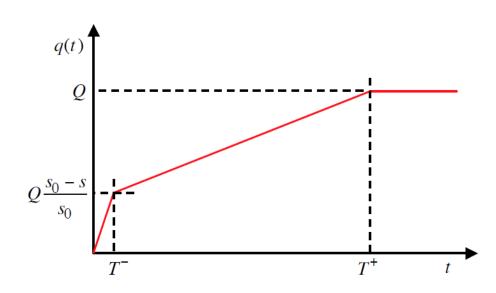
 $w^{-}(w^{+})$: electron (ion) drift velocity $T^{-}(T^{+})$: total electron (ion) drift time (+Q on cathode , -Q on anode)

Charge induced on each electrode by +Q moving through the difference of potential dV:

$$dq = Q\frac{dV}{V_0} = Q\frac{ds}{s_0}$$

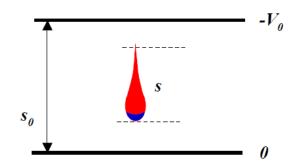
Integrating over s (or time t):

$$q(s) = \frac{Q}{s_0}s$$
 $q(t) = \frac{Q}{s_0}wt$ $i(t) = \frac{dq}{dt} = \frac{Q}{s_0}w$



Charge induction – avalanche multiplication – uniform field

PARALLEL PLATE COUNTERS:



Increase in the number of charges after a path ds:

$$dn = n\alpha ds$$
 $n = n_0 e^{\alpha s}$

Charge induced by electrons: $dq^- = -en_0 e^{\alpha s} \frac{ds}{s_0}$

$$q^{-}(s) = \frac{en_0}{\alpha s_0} (e^{\alpha s} - 1) \approx \frac{en_0}{\alpha s_0} e^{\alpha s} = \frac{en_0}{\alpha s_0} e^{\alpha w^{-}t}$$

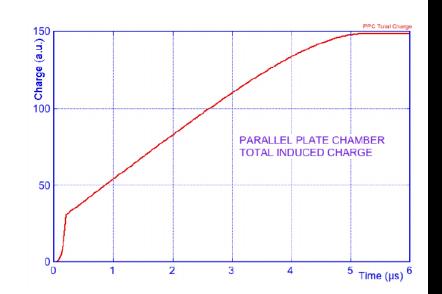
$$i^{-}(t) = \frac{dq^{-}}{dt} = \frac{en_0w^{-}}{s_0}e^{\alpha w^{-}t} = \frac{en_0}{T^{-}}e^{\alpha w^{-}t}$$

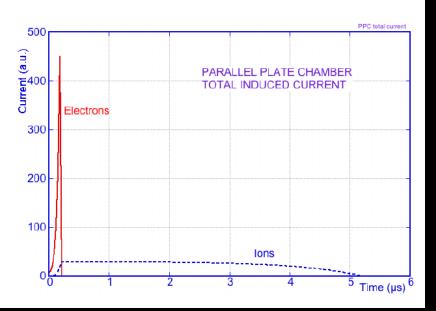
Current signal induced by ions:

$$i^{+}(t) = \frac{en_{0}}{T^{+}} \left(e^{\alpha w^{-}t} - e^{\alpha w^{*}t} \right) \quad 0 \le t \le T^{-}$$

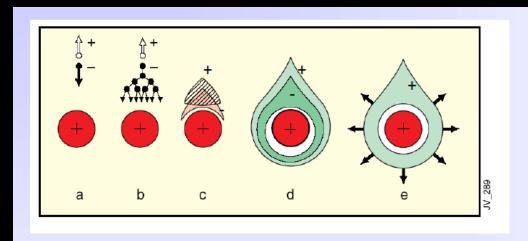
$$i^{+}(t) = \frac{en_{0}}{T^{+}} \left(e^{\alpha s} - e^{\alpha w^{*}t} \right) \quad T^{-} \le t \le T^{+}$$

$$\frac{1}{w^{*}} = \frac{1}{w^{+}} + \frac{1}{w^{-}}$$





SWPC – signal formation



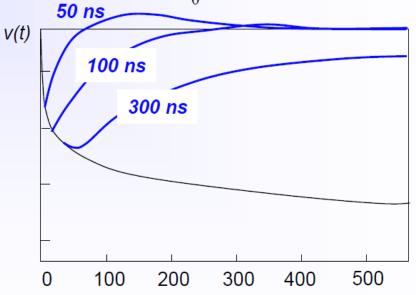
Electrons collected by the anode wire i.e. dr is very small (few μ m). Electrons contribute only very little to detected signal (few %).

lons have to drift back to cathode i.e. dr is large (few mm). Signal duration limited by total ion drift time.

Avalanche formation within a few wire radii and within t < 1 ns.

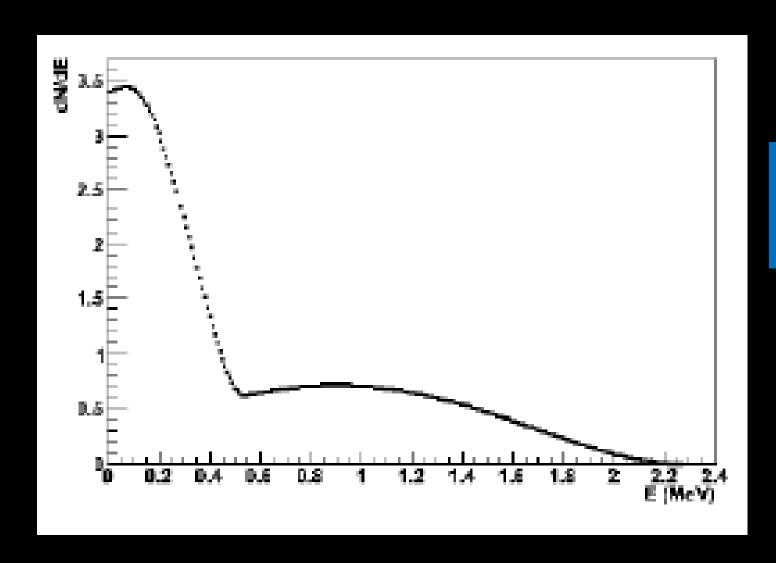
Signal induction both on anode and cathode due to moving charges (both electrons and ions).

$$dv = \frac{Q}{lCV_0} \frac{dV}{dr} dr$$



Need electronic signal differentiation to limit dead time.

Energy spectrum of Sr⁹⁰ radioactive decay



Two decay channels observed:

- 1. β of 546 KeV \rightarrow Y⁹⁰
- 2. β of 2.28 MeV + Zr⁹⁰ stable