PAUL SCHERRER INSTITUT





M. Hildebrandt :: Group Detectors :: Paul Scherrer Institut

Gaseous Detectors and specific characteristics of Drift Chambers

PSI, LTP-Seminar, 08.05.2017



introduction

interaction of ionizing radiation with matter

part I

- fundamental processes in gaseous detectors
- operational modes

part II

- characteristics of drift chambers
- MEG and MEG II drift chambers



Interaction with Matter

Neutral and charged particles are detected through their interactions with matter:

- neutral particles $I(x) = I_0 \cdot e^{-f(x)}$
- charged particles $E(x) = E_0 \cdot e^{-f(x)}$



 charged particles in gases : energy loss by ionization and excitation is described by Bethe-Bloch formula:

$$-\frac{dE}{dx} = 4\pi \cdot N_A \cdot r_e^2 \cdot m_e \cdot c^2 \cdot z^2 \cdot \frac{Z}{A} \cdot \frac{1}{\beta^2} \left[\ln \frac{2m_e \cdot c^2 \cdot \gamma^2 \cdot \beta^2}{I} - \beta^2 - \frac{\delta}{2} \right]$$





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useful constant (and unit) from above:

$$4\pi \cdot N_A \cdot r_e^2 \cdot m_e \cdot c^2 = 0.3071 \frac{MeV}{g/cm^2}$$

→ energy loss
$$-\frac{dE}{dx}$$
 is usually given in units of $\frac{MeV}{g/cm^2}$

energy loss per surface mass density

 \rightarrow energy loss *-dE* along track length *ds*

$$-dE = \frac{dE}{dx} \cdot \rho \cdot ds$$





Interaction with Matter

Neutral and charged particles are detected through their interactions with matter:

- absorption or energy loss changes the properties of incident particle
- density ρ of detection medium influences the change of properties substantially

To follow a charged particle over a long track length, e.g. to measure

- bending radius of the particle trajectory in magnetic field to estimate the particle momentum
- a low-density medium is required:
 - \rightarrow typical application of a gaseous detector

radiation length
$$X_0$$
: $E(x) = E_0 \cdot e^{-\frac{x}{X_0}} \rightarrow E(X_0) = \frac{1}{e}E_0$
 \rightarrow filling gas has long radiation length



Radiation Length

radiation length X₀:

$$E(x) = E_0 \cdot e^{-\frac{x}{X_0}} \quad \Rightarrow \quad E(X_0) = \frac{1}{e}E_0$$

material	Z	A	$X_0[\mathrm{g/cm^2}]$	X_0/ϱ [cm]	$E_{c}[MeV]$
Hydrogen	1	1.01	63	700 000	350
Helium	2	4.00	94	530000	250
Lithium	3	6.94	83	156	180
Carbon	6	12.01	43	18.8	90
Nitrogen	7	14.01	38	30 500	85
Oxygen	8	16.00	34	24000	75
Aluminum	13	26.98	24	8.9	40
Silicon	14	28.09	22	9.4	39
Iron	26	55.85	13.9	1.76	20.7
Copper	29	63.55	12.9	1.43	18.8
Silver	47	109.9	9.3	0.89	11.9
Tungsten	74	183.9	6.8	0.35	8.0
Lead	82	207.2	6.4	0.56	7.40
Air	7.3	14.4	37	30 000	84
SiO_2	11.2	21.7	27	12	57
Water	7.5	14.2	36	36	83

Grupen, 1996



Characteristics of Gaseous Detectors

fundamental processes in gaseous detectors:

- creation of charge
- drift of charge
- amplification of charge
- detection of charge ("real" and induced)



Characteristics of Gaseous Detectors

fundamental processes in gaseous detectors:

- creation of charge
- drift of charge
- amplification of charge
- detection of charge ("real" and induced)

 \rightarrow gaseous electronics



Terminology – Gaseous Electronics

All relevant processes in the filling gas of the detector, in particular

- atomic/molecular physics, e.g. scattering processes
 - rotational, vibrational, electronic excitations in meV – eV range,
 - ionisation, attachment and dissociation
- transport phenomena of charged particles in gases under the influence of electric or electric and magnetic fields, e.g. drift velocity v_d
 - diffusion D_L , D_T
 - $\,\,{}^{_{\rm D}}\,$ magnetic deflection angle $\alpha_{_{\text{Lorentz}}}$

which are explored in

- kinetic theory and transport theory
- discharge physics and low energy plasma physics

are summarised under the expression "gaseous electronics".



Units in Gaseous Electronics

gaseous electronics scales with parameter E/N reason: collision operator in Boltzmann transport equation is proportional to N

particle number density N $N = \frac{n}{1} = \frac{k_B \cdot T}{1}$

 \rightarrow E/N: 1 Townsend = 1 Td = 10⁻²¹ Vm² (Huxley, Crompton, Elford, 1966)

 \rightarrow B/N: 1 Huxley = 1Hx = 10⁻²⁷ Tm³ (Ness, 1991) (Heylen, 1980: 10⁻²³ Tm³)



Sir J.S.E.Townsend 1868-1957



Sir L.G.H.Huxley 1902-1988

<u>@ 20°C, 1 bar:</u>						
1 Td ≈ 250 V/cm						
1Hx ≈ 0.025 T						

following units are often used:

- $[E] = V/cm \rightarrow T$ and p need to be defined
- $[E/p] = V/(cm \cdot torr), V/(cm \cdot atm), V/(cm \cdot mmHg) \rightarrow T$ needs to be defined

• $[E/N] = V \cdot cm^2$

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creation of charge:

 charged and neutral particles dissipate part or all of their energy by generating electron-ion pairs ¬γ: point like

- charged particle: along particle track

primary ionisation n₀:

• $n_0 = \Delta E / w$

• w value: mean energy to produce electron-ion pair

 w value > ionisation potential due to additional excitation of (inner shell) electronic, vibrational or rotational energy levels

separation of charge:

electrical field to avoid recombination and to separate the electron-ion pairs



Gas	Z	Λ	δ	Eex	Ei	I 0	Wi	dE/	dx for MIP	n _p	n _T
			(g/cm ³)		(0	V)		$(MeV/g cm^{-2})$	(keV/cm)	(i.p./cm) ^{a)}	(i.p./cm) ^{a)}
112	2	2	8.38×10^{-5}	10.8	15.9	15.4	37	4.03	0.34	5.2	9.2
He	2	4	1.66×10^{-4}	19.8	24.5	24.6	41	1.94	0.32	5.9	7.8
N ₂	14	28	1.17×10^{-3}	8.1	16.7	15.5	35	1.68	1.96	(10)	56
02	16	32	1.33×10^{-3}	7.9	12.8	12.2	31	1.69	2.26	22	73
Ne	10	20.2	8.39×10^{-4}	16.6	21.5	21.6	36	1.68	1.41	12	39
Ar	18	39.9	1.66×10^{-3}	11.6	15.7	15.8	26	1.47	2.44	29.4	94
Kr	36	83.8	3.49×10^{-3}	10.0	13.9	14.0	24	1.32	4.60	(22)	192
Xe	54	131.3	5.49×10^{-3}	8.4	12.1	12.1	22	1.23	6.76	44	307
002	22	44	1.86×10^{-3}	5.2	13.7	13.7	33	1.62	3.01	(34)	91
a14	10	16	6.70×10^{-4}		15.2	13.1	28	2.21	1.48	16	53。
C4110	34	58	2.42×10^{-3}		10.6	10.8	23	1.86	4.50	(46)	195

a) i.p. = ion pairs

Sauli, 1977



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Sauli, 1977



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a) i.p. = ion pairs

Sauli, 1977

primary ionisation:

- ~100 e⁻/cm for MIP
- not continuously distributed
- clustered along track length



















drift of charge:

- under the influence of external electric or electric and magnetic fields the electrons and positive ions drift towards their corresponding electrode
- electro-negative gas components / impurities will lead to a loss of charge due to electron attachment → loss of signal size
- equilibrium of energy loss in collisions and energy gain between collisions:

 v_{d,electron} ≈ 1-10 cm/μs
 v_{d,ion⁺} ~ 1000-times slower usually kT-limit → constant μ → v_{d,ion} = μ·E

- □ D_T , D_L ≈ 100-500 µm/v1cm
- $\, {}^{_{\rm D}}$ magnetic deflection angle $\alpha_{_{\text{Lorentz}}}$













positive ions: mobility, diffusion (→ Viehland, Mason, McDaniel, Loeb, Townsend)

gas	}. [cm]	$D_{i} = [cm^2/s]$	$(lion \left[\frac{\mathrm{cm/s}}{\mathrm{cm/s}}\right]$	Gas	Ion	Mobility	$[\text{cm}^2 \text{ V}^{-1} \text{ s}^{-1}]$
	Mon[CIII]		Plon [V/cm]	Ar	Ar ⁺	1.54	
H_2	$1.8\cdot 10^{-5}$	0.34	13.0	He	He ⁺	10.4	
He	$2.8\cdot 10^{-5}$	0.26	10.2	CO ₂	CO_2^+	1.09	
Ar	$1.0\cdot 10^{-5}$	0.04	1.7	Ar	CH_4^+	1.87	
O_2	$1.0\cdot 10^{-5}$	0.06	2.2	Ar Ar	$C_2H_6^+$ iC ₄ H ₁₀ ⁺	2.06	
					CO_2^+	1.72	Tavernier, 2010











Neither ~100 e⁻/cm nor ~25 000 e⁻/n-³He-capture is a sufficient amount of charge

 \rightarrow amplification inside gas volume before electronics is needed

amplification of charge:

- exceeds the electrical field ~10 kV/cm the drifting electron (from primary ionisation) gains sufficient energy between collisions so that $\mathcal{E}_{mean} < E_{ionisation}$
 - → possible to create additional electron-ion pairs in collisions with neutral gas atoms or molecules
 - → growing cascade forms an electron avalanche electron multiplication, "gas amplification", gas gain: typically ~10 – 10⁶



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 - → growing cascade forms an electron avalanche electron multiplication, "gas amplification", gas gain: typically ~10 – 10⁶
- avalanche process described by 1st Townsend coefficient, ionisation coefficient:
 - $\ \ \alpha_{\text{Townsend}}$
 - [α] = 1/cm

• total number of electrons: $n = n_0 \cdot e^{\alpha \cdot x}$ or $n = n_0 \cdot e^{\int \alpha(x) dx}$

□ gas gain: n/n₀



ionisation coefficient α



empirical formula (Korff, 1955)

$$\frac{\alpha}{p} = A \cdot e^{-\frac{Bp}{E}}$$

A, B: phenomenological constants

Gas	$A (\text{cm}^{-1}\text{torr})$	$B (\text{V cm}^{-1} \text{torr}^{-1})$
He	3	34
Ne	4	100
Ar	14	180
Xe	26	350
CO ₂	20	466 Sauli, 2014



location of electron avalanche

• wire: diameter $\emptyset \approx 10 - 50 \,\mu\text{m}$, gain $\leq 10^6$



H.Raether, 1949 L.B.Loeb, 1960 G.Charpak, 1968



location of electron avalanche

- wire: diameter $\not{0} \approx 10 50 \ \mu m$, gain $\leq 10^6$
- micro-strips: width ~10 μ m, gain $\leq 10^4$





Oed, NIM A263 (1984) 351

- 1990's: MSGC supposed to be "all-purpose LHC tracking detector"
- today: only D20 at ILL (low gain operation)









location of electron avalanche

- wire: diameter $\emptyset \approx 10 50 \,\mu\text{m}$, gain $\leq 10^6$
- micro-strips: width ~10 μ m, gain $\leq 10^4$
- holes of Gas-Electron-Multiplier (GEM) foil: $\emptyset \approx 50 100 \,\mu\text{m}$, gain $\leq 10^3$
 - ightarrow decoupling of amplification and charge collection



Sauli, NIM A386 (1997) 531

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location of electron avalanche

- wire: diameter $\emptyset \approx 10 50 \,\mu\text{m}$, gain $\leq 10^6$
- micro-strips: width ~10 μ m, gain $\leq 10^4$
- holes of Gas-Electron-Multiplier (GEM) foil: $\emptyset \approx 50-100 \ \mu m$, gain $\leq 10^3$
 - \rightarrow combination of holes and strips: GEM-MSGC



1990s: detector for Inner Tracker HERA-B

Universities Heidelberg, Siegen, Zürich and CERN



location of electron avalanche

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- micro-strips: width ~10 μ m, gain $\leq 10^4$
- holes of Gas-Electron-Multiplier (GEM) foil: $\emptyset \approx 50-100 \ \mu m$, gain $\leq 10^3$
 - ightarrow multi-GEM detectors with dedicated readout structure



- 4-GEM stack
 - (upgrade of ALICE TPC endplate)





location of electron avalanche

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- micro-strips: width ~10 μ m, gain $\leq 10^4$
- holes of Gas-Electron-Multiplier (GEM) foil: $\emptyset \approx 50$ 100 μ m, gain $\leq 10^3$
- parallel-plate gap: MICRO-MEsh GAseous Structure (Micromegas), gain ≤ 10⁵



Giomataris, NIM A376 (1996) 29



Characteristics of Gaseous Detectors

operational modes of gaseous detectors:

- recombination region
- ionization region
- proportional region
- limited proportional region
- Geiger-Müller region
- discharge region




























Recombination Region

region I

- applied voltage (and resulting field) very low
- electrons and ions move with relatively slow speeds
- recombination rate of electrons and ions still considerable
- as voltage (and field) increases, charges move faster
- recombination rate decreases up to point where it becomes 0





Ionization Region

region II

- recombination rate is 0 and no new charge is produced
- the collected charge stays constant despite a change of the applied voltage





Ionization Chamber

ionization chamber placed in proton beam at proton therapy

medical proton cyclotron

- I_{proton} = 1 850 nA
- E_{kin} = 250 MeV
- → huge amount of primary ionisation



dose monitor



position monitor





(True) Proportional Region

region III

- electric field is very strong (> ~10 kV/cm) in certain regions of the detector (e.g. wires, micro-strips, holes of GEM foil, micro-gaps)
- e⁻ from primary ionisation acquire sufficient energy between collisions to produce additional ionisation in e⁻ atom or e⁻ molecule collisions
- the multiplication factor (gas gain) is for a given HV independent of primary ionisation n₀
 - \rightarrow pulse height is proportional to n₀
 - → pulse height is proportional to dissipated energy





Limited Proportional Region

region IV

- free e⁻ are quickly collected, but positive ions move much slower
- each e⁻ avalanche creates a cloud of positive ions which shield the amplification region (e.g. wire) and slowly moves towards the cathode
- this space charge can significantly deteriorate the shape of the electrical field
 - → reduced "effective field" creates less gas amplification
 - → pulse height still increases with increasing number of initial ion pairs but not anymore linearly









Geiger-Müller Region

region V

- space charge created by positive ions becomes completely dominant
- shielding effect of amplification region ¹⁰ increases such that the effective field decreases below the field strength were multiplication starts
 - → self-limited process which will terminate when the same total amount of positive ions have been created
 - → pulse shape and height are
 independent of primary ionisation n₀
 and no longer reflect any properties
 of the incident radiation





Geiger-Müller Region

region V

- depending on amount of "quencher" (photon absorber) in the gas:
 photons emitted from excited atoms and molecules in the avalanche lead to
 - additional avalanches and
 - a spread along the entire length of the wire







Discharge Region

region VI

- continuous breakdown with or without radiation (self-sustaining discharge)
 - → should be avoided to prevent damage to detector





Large Area Gaseous Detectors

- 1908: first wire counter to study natural radioactivity
 E.Rutherford, H.Geiger, 1908
 Geiger, 1913
- 1928: Geiger-Müller counter with single electron sensitivity H.Geiger, W.Müller, 1928 H.Geiger, W.Müller, 1928, 1929
- 1945: proportional tubes
 H.Raether, 1949
- 1968: multi-wire proportional chambers
 G.Charpak, 1968
 G.Charpak *et al.*, 1968





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- 1968: multi-wire proportional chambers
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 G.Charpak *et al.*, 1968
- 1992: Nobel Price in Physics: G.Charpak "...for his invention and development of particle detectors, in particular the multiwire proportional chamber."



G.Charpak (1924 - 2010)



multi-wire proportional chamber (MWPC):

- spatial resolution given by granularity of anode wires (several mm)
- field distortions due to (electrostatic) displacements of anode wires









- characteristic features of drift chamber design:
 - geometry: small variations of E within drift cell
 - [•] gaseous electronics, as $v_d = v_d(E)$: small Δv_d for unavoidable ΔE



elcectrical field without and with potential wire:





elcectrical field without and with potential wire:





elcectrical field without and with potential wire:









elcectrical field without and with potential wire:









requirements for appropriate drift chamber filling gas:

- fast v_d
- *v_d* vs HV plateau
- small diffusion





The drift chamber was "born" in Heidelberg:

- 1968/69: diploma thesis A. H. Walenta
- 1971: A. H. Walenta, J. Heintze and B. Schürlein, Nuclear Instruments and Methods 92 (1971) 373-380 "The multiwire drift chamber - a new type of proportional wire chamber"



A.H.Walenta

J.Heintze

1972: PhD thesis A. H. Walenta: "Lokalisierung von Teilchenspuren durch Messung der Elektronendriftzeiten in grossflächigen Proportionalzählern"

NUCLEAR INSTRUMENTS AND METHODS 92 (1971) 373-380; © NORTH-HOLLAND PUBLISHING CO.

THE MULTIWIRE DRIFT CHAMBER **A NEW TYPE OF PROPORTIONAL WIRE CHAMBER***

A. H. WALENTA, J. HEINTZE and B. SCHÜRLEIN

I. Physikalisches Institut der Universität Heidelberg, Heidelberg, Germany

Received 27 November 1970

In this article a new type of proportional wire chamber is described with large wire distances and accurate position determination by drift time measurement. The electronic circuitry required has been considerably reduced while the space resolution has been improved as compared to a conventional proportional wire chamber. The limit for position accuracy seems to be at present $\sigma = 0.2$ mm.

The complete system with computer read out described in this article has a location accuracy of $\sigma = 0.47$ mm.

 σ = 0.2 mm even with wire pitch ~10 mm







MEG and MEG II experiment

- dedicated experiments for the search for the charge lepton violating decay
 $\mu^{+} \rightarrow e^{+} \gamma$
 - MEG: $B(\mu^+ \rightarrow e^+ \gamma) < 4.2 \cdot 10^{-13} (90\% \text{ CL})$
 - MEG II: x10 improvement in sensitivity
- positron spectrometer (tracker & timing counter) designed for precision measurement of 52.8 MeV/c e⁺
- tracker system
 - high momentum resolution (< 0.4%)
 - high angular resolution (~5mrad)
 - → low-mass tracker for low multiple
 Coulomb scattering contribution
 - \rightarrow very good spatial resolution in sub-mm region
- \rightarrow gaseous detector: drift chamber







MEG experiment: material budget (in units of X_0) along e^+ track

 MEG
 MEG II

 detector module
 2.6·10⁻⁴

 track (7.5 modules)
 2.0·10⁻³

 1.1-1.2·10⁻³







MEG experiment: material budget (in units of X_0) along e^+ track

	MEG	MEG II	300µm Si	50µm Si
detector module	2.6.10-4		32·10 ⁻⁴	5.3·10 ⁻⁴
track (7.5 modules)	2.0·10 ⁻³	1.1-1.2·10 ⁻³	24·10 ⁻³ (?)	4.0·10 ⁻³ (?)







MEG Drift Chamber

- V shaped frame \rightarrow "no" material in sensitive area
- DC module consists of detector two planes
- middle cathode/cathode hood
 12.5 µm polyimid foil, 250 nm Al deposition
- two wire planes: alternating anode and potential wires
- wires planes shifted by d_{anode-pot.wire}







MEG Drift Chamber







MEG Drift Chamber – Characteristics I

field geometry:





potential wires Cu 98/Be 2 Ø 50 μm



anode wires Ni 80/Cr 20 Ø 25 μm



MEG Drift Chamber – Characteristics II

- filling gas He-C₂H₆ (50-50)
 - quite high v_d
 - v_d vs HV plateau
 - long radiation length due to He
 - good HV stability
 due to C₂H₆



"P10": Ar-CH₄ (90-10) "magic gas": Ar-iC₄H₁₀-CHF₃ (75-24.5-0.5)



MEG II Cylindrical Drift Chamber

- based on drift chamber of KLOE experiment at DAΦNE (INFN Frascati)
- single, cylindrical volume (L ~200 cm, Ø_{outer} 60 cm)
- wires with stereo angle and U V views
 → determination of hit position along wire
- small drift cell sizes (7 mm)
- ~1400 anode wires (gold-plated W, Ø 20 μm)
 ~7500 cathode/guard wires (silver-plated Al, Ø 40/50 μm)







L=3.3m, Ø 4m, 52140 wires



MEG II Cylindrical Drift Chamber











MEG II Cylindrical Drift Chamber

drift cell geometry (MEG)



anodecathode


drift cell geometry



anodecathode



drift cell geometry



anode
cathode
guard



drift cell geometry







- filling gas He-iC₄H₁₀ (85-15 or 90-10)
 - Iong radiation length due to high He fraction
 - \rightarrow low multiple scattering, good momentum resolution
 - slow v_d , no v_d vs HV plateau, large diffusion
 - small number of primary ionisation clusters ~13/cm (MEG: He-C₂H₆ 50-50: ~35/cm) → large spacing between clusters





Cluster Counting Technique – in a very tiny nutshell

MEG: He-C₂H₆ (50-50): ~35 e⁻cluster/cm





Cluster Counting Technique – in a very tiny nutshell

MEG: He-C₂H₆ (50-50): ~35 e⁻cluster/cm

in reality





Cluster Counting Technique – in a very tiny nutshell

■ MEG II: He-iC₄H₁₀ (85-15 or 90-10): ~13 e⁻cluster/cm

increased number of "supporting points" along trajectory of incident particle in single drift cell \rightarrow improvement in \Box track fitting accuracy and





MEG Drift Chambers - Performances

	single hit	resolution (σ) momentum	angular
 MEG drift chamber Adam <i>et al.,</i> Eur.Phys.J. C (2013) 73:2365 	r 210 μm z 800 μm	~330 keV (0.6%)	θ 9.4 mrad φ 8.4 mrad
 MEG II drift chamber PhD thesis M.Venturini, 2017 	r ~110 μm (prototype)	~100 keV (0.2%) (MC)	θ ~6 mrad φ ~7 mrad (MC)



Magic Octagon – the Choice of the Filling Gas

quantities in gaseous electronics, like:

- primary ionisation n₀
- drift velocity v_d
- longitudinal, transversal diffusion D_L, D_T
- magnetic deflection angle α_{Lorentz}
- size of electron avalanche (α_{Townsend})
- radiation length X₀
- HV stability, breakdown voltage
- sensitivity to radiation induced degradation
- are a function of: particle number density N, i.e. T and p
 - gas components and mixing ratios
 - electrical and/or magnetic fields
 - rate environment

and the characteristics of the quantities are connected to each other!





- good HV stability
- short radiation length X₀
 - \rightarrow reduced momentum resolution
- bad longterm stability (polymerisation of C-chains)

Vd

D₁,D₁

 α_{Lorentz}

 $lpha_{Townsend}$



Magic Octagon – the Choice of the Filling Gas

quantities in gaseous electronics, like:

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 - rate environment



→ The choice of a counting gas is a complicated task with many compromises and weighing up of advantages, disadvantages and risks and one always has to look at the overall picture!





Wir schaffen Wissen – heute für morgen



Hildebrandt



- 1908: first wire counter to study natural radioactivity
 - E.Rutherford, H.Geiger, Proc. Roy. Soc. A81 (1908) 141-161
 "An Electrical Methode of Counting the Number of α-Particles from Radio-Active Substances"
 H.Geiger, Verh. d. Deutsch. Phys. Ges. 15 (1913) 534-539
 "Über eine einfache Methode zur Zählung von α- und β-Teilchen"
- 1928: Geiger-Müller counter with single electron sensitivity

H.Geiger, W.Müller, Naturwissenschaften 16 (1928) 617-618
"Elektronenzählrohr zur Messung schwächster Aktivitäten"
H.Geiger, W.Müller, Phys. Zeits. 29, 839 (1928) and 30, 489 (1929) and 30, 523 (1929)
"Das Elektronenzählrohr – Wirkungsweise und Herstellung eines Zählrohrs"

• 1945: proportional tubes

H.Raether, Ergeb. Exakt. Naturwiss., Band 22 (1949) 73-120 " Die Elektronenlawine und ihre Entwicklung"

• 1968: multi-wire proportional chambers

G.Charpak, Proc.Int.Symp.Nuclear Electronics, Versailles 10-13 Sep 1968 G.Charpak *et al.*, Nucl. Inst. and Meth. 62 (1968) 262-268 " The use of multiwire proportional counters to select and localize charged particles"



• 1971: drift chamber

A.H.Walenta, J.Heintze, B.Schürlein, Nucl. Inst. and Meth. 92 (1971) 373-380 "The multiwire drift chamber a new type of proportional wire chamber"

■ 1974: Time Projection Chamber

D.R.Nygren, LBL Internal report (Feb 1974); D.R.Nygren PEP 198 (1975)
A.R.Clark *et al.*, "A Proposal for a PEP Facility Based of the Time Projection Chamber"
D.R.Nygren, Physica Scripta Vol.23 (1981) 584-598, "Future Prospects of the TPC idea"

• 1983: GASEOUS DETECTORS FOR PARTICLE PHYSICS: A STILL BLOOMING SOURCE OF NEW INSTRUMENTS

G. CHARPAK CERN, Geneva, Switzerland

Nuclear Instruments and Methods 217 (1983) 5-7 North-Holland Publishing Company

Section I. plenary talks

• 1986: WILL GASEOUS DETECTORS SURVIVE THE RAPID PROGRESS IN THE COMPETING TECHNIQUES?

G. CHARPAK

CERN, Geneva, Switzerland

Nuclear Instruments and Methods in Physics Research A252 (1986) 131-136 North-Holland, Amsterdam



1988: micro-strip gas chamber (MSGC)

A.Oed, Nucl. Instr. and Meth. A263 (1988) 351-359 "Position-sensitive detector with micro-strip anode for electron multiplication with gases"

1990's: huge variety of Micro-Pattern Gaseous detectors (MPGD)

micro-strip gas chamber, gas electron multiplier, micro-mesh-gaseous structure, micro groove detector, micro gap detector, conteur a trous, micro dot avalanche chamber, micro slit gas detector, ...

1996: MICRO-MEsh-GAseous Structure (MICROMEGAS)

Y.Giomataris et al., Nucl. Instr. and Meth. A376 (1996) 29-35 "MICROMEGAS: a high-granularity position-sensitive gaseous detector for high particle-flux environment"

1997: gas electron multiplier (GEM)

F.Sauli, Nucl. Instr. and Meth. A386 (1997), 531-534 "GEM: A new concept for electron amplification in gas detectors"

<u>remark:</u> 1978-1998: Wire Chamber Conference WCC in Vienna since 2001: Vienna Conference on Instrumentation VCI