

Home Search Collections Journals About Contact us My IOPscience

Time projection chambers

This article has been downloaded from IOPscience. Please scroll down to see the full text article.

2010 Rep. Prog. Phys. 73 116201

(http://iopscience.iop.org/0034-4885/73/11/116201)

View the table of contents for this issue, or go to the journal homepage for more

Download details:

IP Address: 137.138.115.224

The article was downloaded on 10/02/2011 at 13:47

Please note that terms and conditions apply.

Time projection chambers

H.J. Hilke

CERN, 1211 Geneva 23, Switzerland

Received 26 November 2009, in final form 14 July 2010 Published 6 October 2010 Online at stacks.iop.org/RoPP/73/116201

Abstract

The time projection chamber (TPC) was originally proposed to permit full reconstruction of events of up to 20 particles at an electron–positron collider. It was intended to provide 3D information for tracking and momentum measurement together with particle identification by multiple ionization sampling, in a compact detector. This powerful combination soon found applications in other fields: at one extreme, for studies of rare events of simple structure; at the other, for heavy ion collisions, handling ever higher particle densities, finally up to several thousand tracks in a single event, still providing information on a track-to-track basis.

Basic physics, performance and limitations of drift chambers in general will be discussed first. Then the characteristics of TPCs will be introduced, as well as their practical realization and performance. The development from the first TPC, 30 years ago, to the present day will be covered and followed by a few final comments on the ongoing studies for future applications.

This article was invited by G Barbiellini.

Contents

Introduction	1	3.3. Performance	22
General properties of DCs	2	4. Particularities of some TPCs	25
2.1. Gas ionization by charged particles	2	4.1. Global aspects	25
2.2. Transport of electrons and ions	5	4.2. Special aspects	27
2.3. Signal formation	10	5. Gas amplification without anode wires	31
2.4. Calibration with laser beams	13	5.1. Parallel plate chamber	31
2.5. Ageing of wire chambers	13	5.2. Micromegas	32
TPC characteristics	15	5.3. Gas electron multiplier	33
3.1. Basic design—PEP4 TPC	15	6. Conclusions	33
3.2. Specific aspects	15	References	34
	General properties of DCs 2.1. Gas ionization by charged particles 2.2. Transport of electrons and ions 2.3. Signal formation 2.4. Calibration with laser beams 2.5. Ageing of wire chambers TPC characteristics 3.1. Basic design—PEP4 TPC	General properties of DCs22.1. Gas ionization by charged particles22.2. Transport of electrons and ions52.3. Signal formation102.4. Calibration with laser beams132.5. Ageing of wire chambers13TPC characteristics153.1. Basic design—PEP4 TPC15	General properties of DCs 2.1. Gas ionization by charged particles 2.2. Transport of electrons and ions 2.3. Signal formation 2.4. Calibration with laser beams 2.5. Ageing of wire chambers 2.6. Ageing of wire chambers 2.7. Basic design—PEP4 TPC 2.8 Particularities of some TPCs 2.9. All Global aspects 3.1. Paccil aspects 3.1. Parallel plate chamber 3.2. Micromegas 3.3. Gas electron multiplier 3.4. Conclusions

1

1. Introduction

The time projection chamber (TPC) is the only electronically read gaseous detector delivering direct three-dimensional track information: for each point on the track, x-, y- and z-coordinates are measured simultaneously. This is of particular importance for pattern recognition in high-multiplicity events. The combination of powerful tracking with particle identification (PID) capacity over a wide momentum range, based on multiple measurements of ionization loss, is the trademark of the TPC. The timely appearance on the market of an affordable multi-sample analogue memory, the charge coupled device (CCD), was an essential ingredient. The major limitation is high event rate.

In 1968, when bubble chambers and spark chambers dominated the field of tracking detectors in high energy physics, multiwire proportional chambers (MWPCs) [1] and almost simultaneously drift chambers (DCs) [2,3] were introduced, with the principal aim of increasing data taking rates. The rapid progress in semiconductor electronics, which had started only a decade before with slow devices, made possible the fast read-out of hundreds and soon thousands of signal channels at affordable cost. This was an essential ingredient to an extremely rapid development of both detector technologies: many different layouts were used and size and complexity of the designs grew constantly. In 1972 already, a MWPC system containing 50 000 wires was under construction. Only 3 years later, the TPC was proposed

by David Nygren [4], a DC providing simultaneously non-projective track recognition, momentum measurement and PID in a compact device.

The basic idea for a detector to be installed around the interaction point of an electron—positron collider was to take a long gas-filled cylinder, with a thin HV electrode in the centre, negatively charged with respect to the MWPCs used for read-out at both ends. The electrons produced along the path of a charged particle traversing the cylinder are drifting to the MWPC and pass through a grid into the multiplication region. They form avalanches near the anode wires. The cathode underneath the wires is segmented into pads and the pulses induced by the avalanches are registered in addition to those from the anode wires. The anode wires are arranged azimuthally around the cylinder axis and their signals provide the radial coordinate. The pads provide the hit coordinate along the wires above them. The third coordinate is obtained from the arrival time of the signals.

Care is taken to produce a homogeneous electric field E along the axis of the cylinder and to align this parallel to the magnetic field B of the solenoid surrounding the TPC, in order to provide an undeformed projection of the original ionization track onto the MPWC plane. A strong magnetic field is necessary for the momentum measurement but is in many cases also useful to improve the point measuring precision due to reduced transverse diffusion. PID is obtained by taking many samples of the track ionization (dE/dx), from the pulse height measurements on the wires or pads.

The first TPC, as part of the PEP4 experiment at SLAC, started taking data in 1983. Rather rapidly, other TPCs were constructed and many followed since, in collider and fixed-target experiments with increasing particle multiplicities. The dimensions grew for some of them significantly, the number of read-out channels even more, encouraged by the fast developments in electronics. This enabled track reconstruction of events with extremely high particle multiplicities, taking full advantage of the unique pattern recognition capabilities. In parallel, the TPC technique found applications in searches for rare and exotic processes with low multiplicities.

We shall start with a discussion of properties applying to DCs in general and then describe particular properties and the performance of TPCs.

The TPC developments until 1992 can be found in a detailed review [5], recent short overviews in [6].

The following terms and numbers will appear repeatedly:

STP = standard temperature and pressure: 0 °C and 1 atm (this is the most commonly used definition of STP, although the new convention of IUPAC demands 1 bar = 1 atm/1.013 25).

Avogadro constant $N_{\rm A}=6.022\times 10^{23}$ molecules per mole. N $\cong 2.68\times 10^{19}\,{\rm mol\,cm^{-3}}={\rm number}$ of molecules per cm³ for an ideal gas at STP; densities of most detector gases at STP are very close to the ideal gas density (e.g. Ar, Ne, CH₄, CO₂). Thermal energy $=3kT/2\cong 0.0353\,{\rm eV}$ at STP.

1 Townsend = 1 Td = 10^{-17} V cm² $\cong 269$ V⁻¹ cm⁻¹ atm⁻¹ at 0 °C and 251 V cm⁻¹ atm⁻¹ at 20 °C (NTP).

Relative speed of a particle $\beta = v/c_0$, where c_0 is the speed of light in vacuum; $\gamma = (1 - \beta^2)^{-1/2}$. A mip = minimum

ionizing particle with $\gamma \sim 3.5$ (the minimum of ionization is rather broad).

Cyclotron frequency $\omega = eB/m \cong 1.76 \times 10^{11} \,\mathrm{s}^{-1}$ for electrons in a magnetic field $B = 1 \,\mathrm{T}$.

2. General properties of DCs

A detailed discussion of all aspects of DCs is presented in [7]. We shall repeatedly refer to this book. A recent account of signal processing also applying to gaseous detectors may be found in [8]. An early overview of wire chamber properties is given in [26].

Powerful simulation programs have been developed and agreement with experimental results has become impressive. The program *Heed* [9] calculates primary ionization produced by fast particles in gases, *Garfield* [10] electric fields, electron and ion trajectories and induced signals and *Magboltz* [11] electron transport properties such as drift velocities and diffusion. These programs are of great help in the planning and optimization of detector setups and operation parameters and for the interpretation of the results.

2.1. Gas ionization by charged particles

2.1.1. Primary clusters. A charged particle crossing the detector loses energy by excitation and ionization of the detector gas, in similar amounts. The ionizing collisions occur randomly, their number k in a segment s along the track following Poisson statistics

$$P(k) = ((s/\lambda)^k/k!) \exp(-s/\lambda), \qquad (2.1)$$

where $\lambda = 1/(N_e \sigma_I)$ is the mean distance between clusters, N_e the electron density of the gas and σ_I the ionization cross-section per electron. The probability to find no cluster in s is $P(0) = \exp(-s/\lambda)$, the probability to find no cluster in s but one in δs is $(\delta s/\lambda) \exp(-s/\lambda)$, i.e. short distances are favoured.

Most ionizing collisions (about 66% in Argon) produce only a single-electron/ion pair. In the other collisions, however, the primary electron receives enough kinetic energy to liberate one or more secondary electrons on other molecules. As the kinetic energies involved are small in most cases, the secondary electrons usually stay close to the first electron/ion pair, forming a 'cluster'. In rare collisions, electrons are ejected with high energy, so-called ' δ -electrons', which leave a detectable ionization trail off the particle trajectory. The average number per centimetre of energetic electrons with energy $E > E_0$ is approximately inversely proportional to E_0 :

$$P(E > E_0) \sim y/(\beta^2 E_0) \text{ cm}^{-1},$$
 (2.2)

with E in keV and y=0.114 for Ar and y=0.064 for Ne [12], $\beta=v/c$. This gives $P=0.011\,\mathrm{cm}^{-1}$ for $\beta=1$ and $E_0=10\,\mathrm{keV}$ in Ar, i.e. one collision with $E>10\,\mathrm{keV}$ in 90 cm on average. The practical range R_p of a 10 keV electron is about 1.4 mm. As R_p is roughly proportional to $E^{1.65}$ in the energy range 1–40 keV, the range for a 1 keV electron is only about 30 μ m.

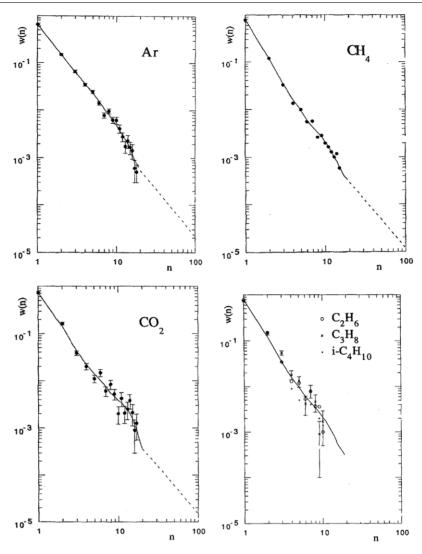


Figure 1. Measurements of cluster size distributions. Full line: eye fit, dashed line: model prediction. (Reprinted with permission from [14]. Copyright 1991, Elsevier.)

Table 1. Measured [14] and theoretical [13] cluster size distributions.

Cluster size	1	2	3	4	5	6	7	8	9	10
Meas. Ar	66	15	6.4	3.5	2.3	1.6	1.1	0.8	0.6	0.5
Meas. CH ₄	79	12	3.4	1.6	1	0.6	0.4	0.3	0.3	0.2
Theor. Ar	80	7.7	2	1.3	0.8	0.6	0.5	0.6	0.8	0.9

Although most of the 'clusters' contain only one electron, clusters with more electrons contribute significantly to the mean total number $n_{\rm T}$ of electrons per centimetre: one finds $n_{\rm T}\gg n_{\rm p}=1/\lambda$, the mean number of clusters per centimetre. For many years there existed only one detailed study providing theoretical estimates for the cluster size distribution [13]. Finally, experimental values have been obtained for cluster sizes up to 18 electrons in Ar, CH₄ and a few other gases, see figure 1 [14]. Table 1 presents some of the experimental and theoretical results.

In Ar one finds on average some 25 clusters cm⁻¹ with a total $n_{\rm T}$ of about 100 electrons cm⁻¹. The most probable value for the total number of electrons is $n_{\rm mp}=42$ electrons cm⁻¹.

The large difference between the mean and the most probable value is caused by the long tail of the cluster distribution. In fact, the largest kinetic energy which can be transmitted to an electron by a heavy particle of mass M and momentum $\gamma\beta cM$ is

$$T_{\text{max}} = 2m_{\text{e}}c^2\beta^2\gamma^2/B; \qquad (2.3)$$

with $B = 1 + (2\gamma m_e/M) + (m_e/M)^2 \to 1 \text{ for } 2\gamma m_e/M \ll 1.$

It follows that a 1 GeV proton can transmit a maximum of about 1 MeV to an electron, but with a probability of only about 10^{-5} cm⁻¹ in Ar.

Both the Poisson distribution of the clusters along the particle track and the cluster size distribution influence the spatial resolution in a DC, whether it is determined by the arrival time of the first n electrons or by centre-of-gravity calculation from induced pulses. In addition, energetic δ -electrons can produce signals significantly off the track.

2.1.2. Total ionization—dE/dx. Over a wide energy range, elementary particles heavier than electrons lose energy predominantly by atomic excitation and ionization, whilst

Table 2. Properties of gases at 20 °C, 1 atm. n_p , n_T : mean primary and total number of electron–ion pairs per centimetre; w: average energy dissipated per ion pair; E_1 , E_x : lowest ionization and excitation energies [15, p 292].

Gas	$n_{ m p}$	n_{T}	w (eV)	<i>E</i> _I (eV)	$E_{\rm x}$ (eV)	$\begin{array}{c} \rho \\ (\mathrm{mgcm^{-3}}) \end{array}$
Не	4.8	7.8	45	24.5	19.8	0.166
Ne	13	50	30	21.6	16.7	0.84
Ar	25	100	26	15.7	11.6	1.66
Xe	41	312	22	12.1	8.4	5.50
CH_4	37	54	30	12.6	8.8	0.67
C_2H_6	48	112	26	11.5	8.2	1.26
i-butane	90	220	26	10.6	6.5	2.49
CO_2	35	100	34	13.8	7.0	1.84
CF_4	63	120	54	16.0	10.0	3.78

for electrons bremsstrahlung becomes dominant already at relatively low energies. The mean energy loss for heavy particles of charge z in detectors is well described by the modified Bethe–Bloch equation [15, p 268]:

$$\langle dE/dx \rangle = Kz^2 (Z/A) (1/\beta^2) [0.5 \ln(2m_e c^2 \beta^2 \gamma^2 T_{\text{max}}/I^2) - \beta^2 - \delta/2],$$
 (2.4)

where $K = 4\pi N_A r_e 2m_e c_0^2$, $N_A = \text{Avogadro constant}$, m_e , r_e are the mass and classical radius of the electron and I is the mean excitation energy of the atom.

The energy loss is a function only of the particle velocity β , except for a minor dependence on the particle mass through the maximum energy transfer T_{max} . A minimum ionizing particle (mip) has $\beta \gamma = 3$ –4 and a minimum energy loss per centimetre of 1–2 MeV g⁻¹ cm² for all materials, except for hydrogen with 4 MeV g⁻¹ cm² loss. The minimum is followed at higher energies by a logarithmic rise which tends to level off faster for denser materials as described by the density correction δ (being a function of $\beta \gamma$).

The translation into ionization is obtained by introducing w, the mean energy lost to produce an electron/ion pair: $w = \langle dE/dx \rangle/n_T$.

It turns out that w is rather independent of particle energy, except for the lowest energies (<100 eV for electrons, <10–30 keV for protons and below some MeV for α -particles) [16]. In rare gases (except possibly for He) the values are also very close for different particles, from electrons to α -particles; for some organic vapours they are up to 15% higher for α -particles. For rare gases one finds w=(1.4-1.8) $E_{\rm I}$ and for common molecular gases w=(2.1-2.5) $E_{\rm I}$, see table 2. The difference to the ionization energy $E_{\rm I}$ is the result of losses due to excitation and to slow electrons with kinetic energies below the first excitation level.

In a gas mixture with one component A having an excitation level higher than the ionization level of component B, excited molecules of A often induce significant additional ionization of B molecules, as observed, e.g., in He and Ne even with minute impurities: the addition of 0.13% Ar to He changed w from 41.3 to 29.7 eV per ion pair. This energy transfer is called the *Jesse effect* or *Penning effect*, when metastable states are involved.

The Bethe–Bloch equation describes the totality of losses, including those of δ -electrons with energies up to the maximum

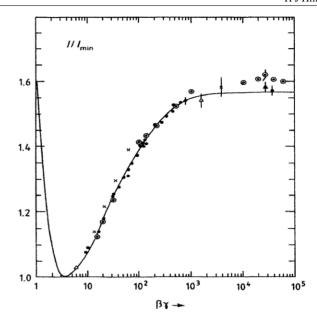


Figure 2. Relativistic rise of ionization I, normalized to the minimum I_0 , as a function of $p/mc = \beta \gamma$. The curve is from simulation [18], the crosses are measurements. (Reprinted with permission from [18]. Copyright 1984, Elsevier.)

possible energy $T_{\rm max}$. These electrons may leave the sensitive zone of a detector element. As one is interested in the ionization near to the particle trajectory, it is sensible to replace $T_{\rm max}$ by a cut-off energy $T_{\rm cut}$. This leads to the formula for the mean restricted energy loss for $T_{\rm cut} \ll T_{\rm max}$ [15, p 270].

$$\langle dE/dx_{\text{restricted}} \rangle = Kz^{2}(Z/A)(1/\beta^{2})$$

 $\times [0.5 \ln(2m_{e}c^{2}\beta^{2}\gamma^{2}T_{\text{cut}}/I^{2}) - \beta^{2}/2 - \delta/2].$ (2.5)

The cut-off obviously depends on the read-out technique and could lie around 10-50 keV outside a magnetic field and higher inside. Assuming w is independent of the particle energy for a given gas, the *ionization measured* in the detector should follow the restricted energy loss. That means, that for a sample of length L along the particle trajectory, the mean total number of electrons deposited is

$$\langle n(L) \rangle = L \langle dE/dx_{\text{restricted}} \rangle / w = n_{\text{T}}L.$$
 (2.6)

Figure 2 shows a measured relativistic rise and the plateau [17] in Ar/CH₄ (90/10) compared with simulation [18] and figure 3 the density effect for various gas pressures [19]. The lowering of the Fermi-plateau for higher pressure is significant.

The distribution of n(L) for small samples L is very broad, with a long tail towards the high energies, caused by energetic δ -rays, see figure 4. The mean and variance of this distribution are, therefore, ill-defined and it is better to characterize the pulse height distribution of the sample by its *most probable number of electrons per centimetre*, $n_{\rm mp}$, and its full-width-half-mean, FWHM. To obtain a reasonable accuracy on $n_{\rm mp}$ for PID by dE/dx, many samples are required, see section 3.3.2.

Table 2 shows experimental values for some of the common detector gases for a minimum ionizing particle (mip). Results from different measurements vary up to a few per cent for $n_{\rm T}$, w and in particular for $n_{\rm p}$, as cluster separation is experimentally difficult.

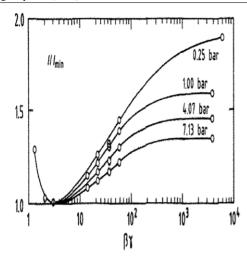


Figure 3. Density dependence of the relativistic rise in Ar/CH₄ (90/10). (Reprinted with permission from [19]. Copyright 1979, Elsevier.)

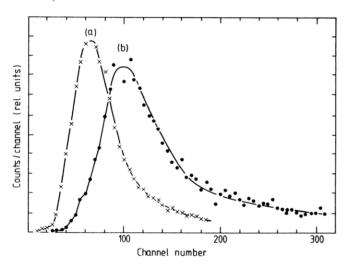


Figure 4. Measured pulse height distribution for a 2.3 cm sample in Ar/CH₄ at 1 atm: (a) 3 GeV/c protons, (b) 2 GeV/c electrons. (Reprinted with permission from [19]. Copyright 1979, Elsevier.)

2.1.3. Effective number of electrons $N_{\rm eff}$. Due to the electron clustering, for several quantities of interest to us, the number of electrons relevant is the number $N_{\rm eff}$ of independently acting electrons and not the total number. We follow some of the arguments of [7].

Take the simple case of a track segment L and determine the precision of the centre of gravity of the charge deposition, assuming m pointlike clusters with n_i electrons are deposited at x_i . The centre of gravity is

$$X_{\rm cg} = \sum (x_i n_i) / \sum n_i = \sum (x_i n_i) / N.$$
 (2.7)

The sum goes from 1 to m, N is the total number of electrons in L. Let x = 0 be the centre of L. The probability distribution for each x_i is

$$F(x_i) dx_i = (1/L) dx_i.$$
 (2.8)

For every fixed n_i the average is $\langle x_{cg} \rangle = 0$ and

$$\langle x_{\rm cg}^2 \rangle = \left\langle \sum_i x_i^2 n_i^2 \right\rangle / N^2 = (L^2/12) \left(\sum_i n_i^2 \right) / N^2$$

= $(L^2/12)/N_{\rm eff}$, (2.9)

as definition of $N_{\rm eff}$ with

$$N_{\rm eff} = N^2 / \left(\sum n_i^2\right). \tag{2.10}$$

 $N_{\rm eff}$ is always $\leq m$. Take as an extreme case m=2 clusters with $n_2 \gg n_1$. Then $N_{\rm eff}$ is close to 1, as the effect of the small cluster is negligible. For the cluster distribution calculated by [13], approximately $N_{\rm eff}=m^{0.54}$ has been obtained [7].

The values of $N_{\rm eff}$ are thus very small for situations with well separated clusters. This changes when clusters start mixing through diffusion. In the limit of very large diffusion, $N_{\rm eff}$ will approach the total number N of electrons. This declustering through diffusion has been observed [20] and simulated [7]. For a mean distance between clusters of 1/m = 0.37 mm, the cluster distribution of [13] and a sample length of 4 mm, the simulation indicates $N_{\rm eff} = 0.3$ times m without diffusion, equal to m for rms diffusion of 1.6 mm and twice m for diffusion of 4 mm.

A similar effect is caused by the fluctuations in the gas amplification and in practice these two effects are combined. The combined action on the spatial resolution from centroid determination in the TPC has been analysed in detail and compared with measurements [21].

2.2. Transport of electrons and ions

In this section, only the main results can be presented, often without derivation. A detailed account of DC properties is given in [7], to which we shall refer at several occasions.

Without external fields, electrons and ions will be in thermal equilibrium and their movement will be described by a mean instantaneous velocity v given by $(m/2)v^2 = (3/2)kT$ and an isotropic diffusion coefficient D. When an electric field E is applied, electrons and ions will acquire drift velocities u antiparallel, respectively, parallel to E, with u much smaller than v. In the presence of an additional magnetic field, the drift directions will be determined by both E and B. Diffusion will no more be isotropic but will show a coefficient for longitudinal diffusion D_L , along E, which may be quite different from D_T , describing diffusion at right angles to E. The big mass difference between electrons and ions results in very different behaviour.

In the following sections, a simple model will bring out the main characteristics and explain qualitatively the results from measurements. This model is based on the approximation of a single instantaneous velocity v and a single *effective cross-section* $\sigma(v)$ describing the elastic and inelastic spectrum. More rigorous treatments including the distribution of v around its mean value achieve quantitative agreement with measurement for many cases.

2.2.1. Drift velocities.

Drift of electrons. When moving in a gas, electrons—because of their light mass—scatter isotropically and forget any preferential direction after each collision with an atom. If an electric field E is present in the drift volume, an electron will be accelerated between collisions and acquire a drift velocity u,

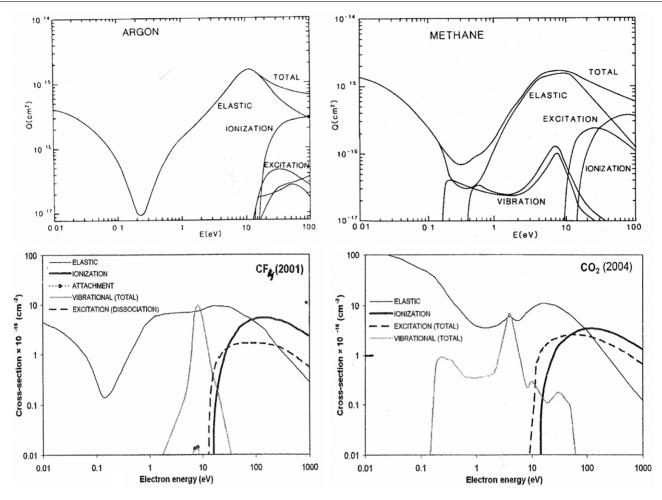


Figure 5. Electron collision cross-sections used in Magboltz [11]. (Reprinted with permission from [23]. Copyright 1989, Elsevier.)

which is equal to the acceleration times the average time τ since the last collision, τ being also the mean time between collisions:

$$u = eE\tau/m. (2.11)$$

The *mobility* μ , defined by $u = \mu E$, is not constant, as τ depends on E.

In the next collision, the electron will lose a fraction Δ of the energy ε_E picked up between collisions and over a distance x there will be a balance between the energy lost and the energy gained:

$$(x/u)(1/\tau)\Delta\varepsilon_E = eEx, \qquad (2.12)$$

as $(x/u)(1/\tau)$ is the number of collisions over a distance x.

The mean time τ between collisions is related to the collision cross-section σ per molecule, the number density N of gas molecules and the *mean instantaneous velocity* v by

$$(1/\tau) = N\sigma v. \tag{2.13}$$

Including the thermal energy, the total energy ε of the electron is given by

$$\varepsilon = \varepsilon_E + (3/2)kT = (m/2)v^2. \tag{2.14}$$

In particle detectors, the approximation $\varepsilon \gg (3/2)kT$ for drifting electrons is fulfilled for many gases; these gases—such as Ar and CH₄—are called 'hot' gases in contrast to 'cold'

gases as CO_2 with ε close to (3/2)kT. One then obtains

$$u^{2} = (eE/mN\sigma)\sqrt{(\Delta/2)}$$
 (2.15)

and

$$v^2 = (eE/mN\sigma)\sqrt{(2/\Delta)}$$
 for $\varepsilon \sim \varepsilon_E \gg (3/2)kT$. (2.16)

A more rigorous theory assuming a Druyvesteyn distribution for the random velocities v adds a multiplication factor of about 0.85 to the right sides [7]. To achieve accuracies of a per cent or better, it is necessary to go one step further and to apply a multiterm $(n \ge 4)$ spherical harmonic expansion treatment of the Boltzmann equation [22] or use Monte Carlo, as done in the present Magboltz 2 [11].

As can be seen, E and N only appear as E/N, the reduced electric field. A special unit, the Townsend (Td), is often used for E/N: $1 \text{ Td} = 10^{-17} \text{ V cm}^2$.

Both σ and Δ depend on ε . For energies ε below the first excitation level, the scattering is elastic and $\Delta \sim 2m/M \sim 10^{-4}$ for molecules of mass M. Low σ and large Δ are required for high drift speeds. An example of a low σ is the so-called 'Ramsauer dip' in the cross-section σ observed in heavy rare gases and in CH₄, see figure 5 [23], which leads to high drift velocities in Ar–CH₄ mixtures at low E-values. This was an important reason for the choice of these gases in several TPCs.

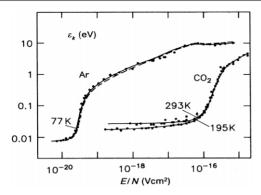


Figure 6. Characteristic electron energy ε_k derived from measurements of diffusion and drift velocity, as a function of reduced electric field E/N for a 'hot' and a 'cold' gas, Ar and CO₂, respectively [7, 25(a)]. 10^{-17} V cm² = 251 V cm⁻¹ atm⁻¹ at 20 °C. (Reprinted with permission from [7]. Copyright 2008, Springer Science + Business Media.)

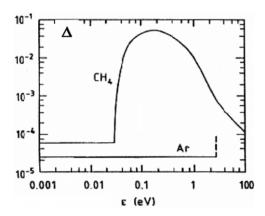


Figure 7. The fraction Δ of energy lost per collision as a function of mean energy ε of the electron [24].

For gas mixtures with number densities n_i $(N = \sum n_i)$, the effective σ and Δ are given by

$$\sigma = \sum n_i \sigma_i / N \tag{2.17}$$

and

$$\Delta \sigma = \sum n_i \Delta_i \sigma_i / N. \tag{2.18}$$

For some gases, σ and Δ have been calculated from precise measurements of drift velocity u (to 1%) and diffusion D/μ (to 3–5%), see, e.g., [24]. Figure 6 shows the *characteristic energy* $\varepsilon_k = (2/3) \varepsilon$ derived in another study [25] in the same way for two extreme cases, 'cold' CO_2 and a 'hot' Ar. In 'cold' gases electrons lose energy to low vibrational and rotational levels, which keeps ε low. Figure 7 presents calculated values for Δ as a function of ε [24].

Drift velocities for different gas mixtures show very different values and dependence on the electric field. At low fields, they in general rise with electric field. Then for many gases they go through a more or less pronounced maximum and minimum before rising again. Drift velocities are shown in figure 8 for a few gases [15, 26]. For a large number of gas mixtures, velocities calculated with Magboltz 2 version 7.1 can be found in [33].

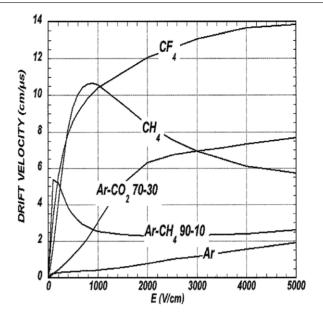


Figure 8. Computed drift velocities of electrons in a few gases. (Reprinted with permission from [15]. Copyright 2008, The Regents of the University of California.)

Drift of ions. A detailed account of mobility and diffusion of ions in gases is given in [27]. Because of their heavy mass m_i , ions lose a significant fraction Δ of their energy in a collision with a gas molecule M (on average $\Delta = 2m_i M/(m_i + M)^2$), so that their random energy will stay close to thermal energy and the momentum will only be partially randomized. As a result, diffusion will be much smaller than for electrons and the mobility will stay constant up to high fields E.

For particle detectors, the low E-field approximation is valid, except in the regions of high gas amplification. The random velocity is taken as thermal, i.e. the relative velocity $v_{\rm rel}$ between the ion and the gas molecules of masses $m_{\rm i}$ and M, respectively, is assumed to be

$$v_{\rm rel}^2 = v_{\rm ion}^2 + v_{\rm eas}^2 = 3kT(m_{\rm i}^{-1} + M^{-1}).$$
 (2.19)

The average time between collisions is $\tau = 1/(N\sigma v_{\rm rel})$ and an argumentation similar to the one followed for electrons [7] leads to

$$u = (m_i^{-1} + M^{-1})^{1/2} (3kT)^{-1/2} eE/(N\sigma).$$
 (2.20)

At low *E*-fields, the drift velocities are proportional the E/N and the mobilities are constant, as σ is practically constant in the energy spectrum of the ions. A typical value is $u \sim 2 \,\mathrm{m \, s^{-1}}$ for CH₄ at $E = 100 \,\mathrm{V \, cm^{-1} \, atm^{-1}}$, to be compared with a thermal velocity around 650 m s⁻¹ at STP.

For very high E-fields, the drift velocities are proportional to the square root of E/N or of E/p, if temperature is kept constant. Figure 9 shows measured values for E/p in noble gases: both extremes are visible, with a transition for Ar between 25 and 150 kV cm⁻¹ atm⁻¹.

In *gas mixtures* it is expected that after a short distance of the order of a centimetre only the species with the lowest ionization energy is left over from the various ion types produced originally, because the *charge transfer* cross-sections are big and similar to other ion–molecule scattering cross-sections [119]. Measurements in binary and ternary

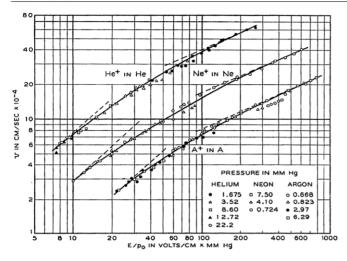


Figure 9. Drift velocities of singly charged ions of noble gases. (Reprinted with permission from [28]. Copyright 1957, American Physical Society.)

mixtures support this assumption, showing a simple relation (Blanc's law) for the ion mobility μ :

$$1/\mu = \sum p_k/\mu_k,\tag{2.21}$$

where p_k are the volume concentrations of the components and μ_k the mobilities in the component k [25]. This relation is expected, if the type of drifting ions does not change.

In an argon–isobutane–methylal mixture the charge exchange from isobutane to methylal becomes inefficient only for concentrations of methylal below about 3%.

Magnetic field effects. Langevin has derived a relation providing a good approximation for many practical cases:

$$u = (e/m)\tau |E|(1/(1+\omega^2\tau^2))\{E^* + \omega\tau [E^* \times B^*] + \omega^2\tau^2 (E^* \cdot B^*)B^*\},$$
(2.22)

where m, e and u are the particle's mass, charge and velocity vector; E and B are the electric and magnetic field vectors $(E^*$ and B^* are unit vectors); $\omega = (e/m)|B|$ and carries the sign of e; τ is the *mean time between collisions*.

For ions, $\omega \tau \sim 10^{-4}$. Therefore, magnetic fields have negligible effect on ion drift.

For electrons, u is along E, if B = 0, with

$$\mathbf{u} = (e/m)\tau \mathbf{E},\tag{2.23}$$

the same relation which was derived in section 2.2.1.1.

For large $\omega \tau$, i.e. large B, u tends to be along B, but if $E \cdot B = 0$, large $\omega \tau$ turns u in the direction of $E \times B$. In DCs, two cases are of particular interest.

E orthogonal to B. With $E \cdot B = 0$ and choosing $E = (E_x, 0, 0)$ and $B = (0, 0, B_z)$, we get

$$u_x = (e/m)\tau |E|/(1+\omega^2\tau^2),$$
 (2.24)

$$u_v = -(e/m)\tau \ \omega \tau |E|/(1 + \omega^2 \tau^2),$$
 (2.25)

$$u_{z} = 0 \tag{2.26}$$

and

$$tg\psi = u_y/u_x = -\omega\tau. \tag{2.27}$$

From a measurement of this *Lorentz angle* ψ , τ can be obtained. As ψ leads to an increased spread of arrival times and sometimes also increases the lateral spread, a small $\omega \tau$ would help, but momentum resolution usually requires a strong B.

The absolute value of u is

$$|\mathbf{u}| = (e/m)\tau |\mathbf{E}| (1 + \omega^2 \tau^2)^{-1/2} = (e/m)\tau |\mathbf{E}| \cos \psi.$$
 (2.28)

The component of E along u determines the drift velocity (Tonks' theorem), independent of the drift direction. This is well verified by experiments.

E nearly parallel to B. This is the case in the TPC. Assuming *E* along *z* and the components $|B_x|$ and $|B_y| \ll |B_z|$, one finds in first order

$$u_x/u_z = (-\omega \tau B_y/B_z + \omega^2 \tau^2 B_x/B_z)/(1 + \omega^2 \tau^2),$$
 (2.29)

$$u_v/u_z = (\omega \tau B_x/B_z + \omega^2 \tau^2 B_v/B_z)/(1 + \omega^2 \tau^2).$$
 (2.30)

In a TPC this will produce a displacement after a drift length L of $\delta_x = Lu_x/u_z$ and $\delta_y = Lu_y/u_z$. As example, let us assume $B_x = 0$ and $\omega \tau \gg 1$. Then $u_y/u_z \sim B_y/B_z$. A good alignment of the E and B axes is therefore important to keep necessary corrections low. From measurements with both B-field polarities and different field strengths, the remaining B_x , B_y can be determined, as well as τ .

If B_x and B_y are small with respect to B_z , u_z remains unaffected by B.

2.2.2. Diffusion. For a long time, diffusion in gases without magnetic fields has been assumed to be isotropic. In 1967, however, it was found experimentally [29] that the longitudinal diffusion $D_{\rm L}$ for electrons along E can be significantly different from the transversal diffusion $D_{\rm T}$. Subsequently it has been established that this is usually the case for electrons [33].

To achieve high precision in calculations of diffusion coefficients, the use of Monte Carlo techniques [11] or a multiterm expansion treatment of the Boltzmann equation [22] is even more important than for calculations of drift velocities. Errors introduced by a 2-term treatment were found to be around 5% for electron diffusion in nitrogen.

Isotropic diffusion. The simpler case of isotropic diffusion shall be treated first. It describes well the behaviour of ions, except for very large electric fields E, and is a reasonable approximation for electrons in some cases.

Due to the random collisions with the gas molecules, a pointlike cloud of electrons or ions starting at t=0 to drift in the *z*-direction due to *E* will at time *t* show a Gaussian density distribution

$$N = (4\pi Dt)^{-3/2} \exp(-r^2/(4Dt)), \qquad (2.31)$$

with $r^2 = x^2 + y^2 + (z - ut)^2$, D being the diffusion coefficient. In any direction from the cloud centre, the mean squared deviation of the electrons is

$$\sigma_x = (2Dt)^{1/2}. (2.32)$$

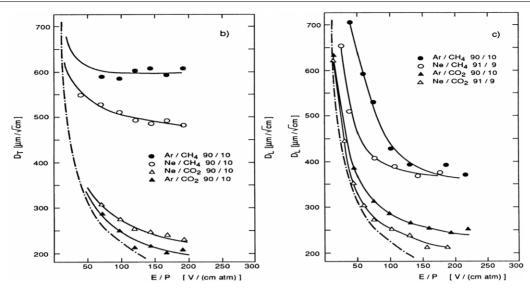


Figure 10. Transverse $(D_{\rm L}^*)$ and longitudinal $(D_{\rm L}^*)$ diffusion constants for electrons, the rms spread after 1 cm drift. The dashed–dotted line indicates the thermal limit [31].

D is usually determined from the variation of the lateral cloud width as a function of z = tu

$$\sigma_{\rm x}^2 = 2Dz/u. \tag{2.33}$$

Related to the *diffusion coefficient D* is the *diffusion constant* D^* , with

$$D^{*2} = 2D/u, (2.34)$$

leading to

$$\sigma_{\rm r} = D^* z^{1/2}. (2.35)$$

The diffusion constant D^* is the diffusion spread for a single electron after 1 cm drift. For a drift length L cm the spread will be $\sigma_x = D^* \sqrt{L}$.

In terms of the microscopic picture, D is given by

$$D = \lambda^2/(3\tau) = v\lambda/3 = v^2\tau/3 = (2/3)(\varepsilon/m)\tau, \quad (2.36)$$

with λ being the mean free path, $\lambda = v\tau$ and ε the mean energy. As $u = (eE/m)\tau = \mu E$, with u = drift velocity and $\mu =$ mobility,

$$\varepsilon = (3/2)(D/\mu)e. \tag{2.37}$$

The mean energy ε can be determined by a measurement of D and u.

Instead of ε , one often finds the 'characteristic energy'

$$\varepsilon_k = (2/3)\varepsilon = eD/\mu.$$
 (2.38)

The lateral diffusion width σ_x after a drift distance L can also be expressed in terms of the mean energy ε :

$$\sigma_{\rm x}^2 = 2Dt = 2DL/(\mu E) = (4/3)\varepsilon L/(eE).$$
 (2.39)

To obtain low diffusion and thus good spatial resolution in DCs, a low electron energy and high electric fields are required. The lower limit for ε is the thermal energy $\varepsilon_{th} = (3/2)kT$. In this limit, the relationship known as *Einstein or Nernst–Townsend formula* follows:

$$D/\mu = kT/e. \tag{2.40}$$

The minimum diffusion width is thus

$$\sigma_{x,\min}^2 = (kT/e)(2L/E).$$
 (2.41)

As can be seen in figure 10, this minimum is approached for electrons in 'cold gases' such as Ar/CO_2 up to $E \sim 100 \, V \, cm^{-1}$ at 1 atm, for 'hot gases' such as Ar/CH_4 only for much lower fields. Ions follow this thermal limit always, except in very high fields.

Anisotropic diffusion. For anisotropic diffusion in an electric field E along z, the transversal diffusion coefficient D_T is defined by

$$\sigma_{\rm r}^2 = 2D_{\rm T}t = 2D_{\rm T}L/u_z \tag{2.42}$$

and the longitudinal diffusion coefficient D_L along E is by

$$\sigma_{\rm t}^2 = u^{-2}\sigma_{\rm r}^2 = 2D_{\rm L}u_{\rm r}^{-2}t = 2D_{\rm L}L/u_{\rm r}^3.$$
 (2.43)

Measurements of lateral diffusion, drift velocity and spread of arrival times σ_t allow determination of both diffusion coefficients. To obtain precise results, great care has to be taken to measure single electrons with good efficiency [30].

Magnetic field effects. A magnetic field B along z will curl up electrons in the x-y projection between collisions. The lateral diffusion will thus be reduced:

$$D_{\rm T}(B)/D_{\rm T}(0) = 1/(1 + \omega^2 \tau^2).$$
 (2.44)

This reduction is essential for most TPCs operating at 1 atm with their long drift distances. At higher pressure, the shortening of τ reduces the gain.

A more rigorous treatment of averages [7] shows that at high *B* the ratio is changed to

$$D_{\rm T}(B)/D_{\rm T}(0) = 1/(C + \omega^2 \tau_2^2)$$
 with a different τ . (2.45)

This behaviour was indeed verified [32] by measuring D(B) over a wide range of B [24]. In an Ar/CH₄ (91/9) mixture the data could be fitted with $\tau = 40$ ps, $\tau_2 = 27$ ps and

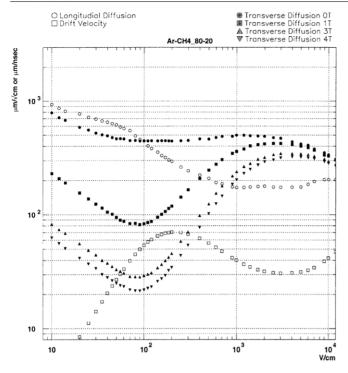


Figure 11. Transverse and longitudinal diffusion constants for electrons in Ar/CH₄ (80/20), in μ m/cm^{1/2}, as a function of electric and magnetic fields. Drift velocity in μ m ns⁻¹ [33(b)].

C = 2.8. The low field behaviour is followed up to about 2 kG, the high field behaviour is approached above about 3.5 kG.

The *longitudinal diffusion* remains unchanged : $D_L(B) = D_L(0)$.

Figures 10-12 show transverse and longitudinal diffusion for a few DC gases. A database for many gas mixtures, calculated with Magboltz 2, version 7.1, has been compiled in [33]. It contains diffusion coefficients as a function of E and B (for transverse diffusion only) and drift velocities as a function of E. One example is shown in figure 11.

2.2.3. Electron attachment. Drifting electrons may be absorbed by electronegative components or impurities in the gas mixture. Halogenides (e.g. CF₄) and oxygen have particularly strong electron affinities. Two-body and three-body attachment processes are distinguished [34].

In the *two-body process*, the molecule may or may not be broken up:

$$e^- + AX \to AX^{-*} \to A(\text{or } A^*) + X^-(\text{or } X^{-*})$$
 (2.46)

or

$$e^- + AX \rightarrow AX^{-*} \rightarrow AX^- + \text{energy}.$$
 (2.47)

The attachment rate R is proportional to the density N:

$$R = c\sigma N \tag{2.48}$$

for an electron velocity c and attachment cross-section σ . The rate constants of freons and many other halogen-containing compounds are known [35].

The Bloch–Bradbury process is the best known three-body process [36]: an electron is attached to a molecule through

the stabilizing action of another molecule. This process is important in O_2 for the attachment of electrons with energies below 1 eV, forming an excited unstable state with a lifetime τ of the order of 10^{-10} s. A stable ion will be formed only if the excitation energy is carried away during τ by another molecule. As the attachment rate depends on the product of the concentrations of oxygen and of the stabilizing molecules, it is proportional to the square of the gas pressure [7]:

$$R = \tau c_e c_2 \sigma_1 \sigma_2 N(O_2) N(X).$$
 (2.49)

Here c_e is the electron velocity, c_2 the relative thermal velocity between O_2 and X. An O_2 contamination of 1 ppm in an Ar/CH₄ (80/20) mixture at 8.5 atm causes an electron absorption of 3% m⁻¹ at a drift speed of 6 cm μ s⁻¹.

2.3. Signal formation

All TPCs operated in experiments have so far used MWPCs for the read-out. Presently, for the first time, a TPC with a different read-out is installed in an experiment at T2K (see section 3.6.2). We shall, therefore, first discuss the MWPCs. The signal is determined by induction from the moving charges, the gas amplification near the anode wire and the response of the electronics. The shaping times in the MWPC have to be between 100 and 200 ns typically. This results in a noise of at best some 300 but often >1000 electrons. In addition, only a fraction of the total ionization is normally available with this pulse shaping (see section 2.3.1) For this reason, a gas amplification of >10000 is usually required. This amplification is obtained by avalanche multiplication of the original electron in the high electric field of up to $E \sim$ $200 \,\mathrm{kV} \,\mathrm{cm}^{-1}$ around the thin anode wire (diameter $\sim 20 \,\mu\mathrm{m}$) at 1 atm.

2.3.1. Signal induction. We shall consider the simple case of a single anode wire on the axis of a cylindrical cathode tube. This describes well the general situation in a MWPC, because inside about 1 mm around the anode wires the field is radial and almost all electrons are produced closer than $100 \, \mu \text{m}$ to the wire [37].

Signals are always produced by induction from the moving charges.

In any electrode configuration, the current I_R induced on the read-out electrode R is given by

$$I_R = -q\mathbf{E}_{\mathbf{W}} \cdot \mathbf{v},\tag{2.50}$$

where q is the signed charge moving with the vectorial velocity v and $E_{\rm W}$ is a vectorial 'weighting field', a conceptual field defined by applying +1 V on R and 0 V on all other electrodes. The unit of $E_{\rm W}$ is 1 cm⁻¹ [38, 39].

The actual v is calculated by applying the normal operation voltages, including possibly a B-field.

In the special case of a two electrode system like the wire tube or a parallel plate chamber, $E_{\rm W}=E_{\rm op}/V$, where $E_{\rm op}$ is the actual operating field obtained with the voltage V on R (the anode wire) and zero V on the cathode.

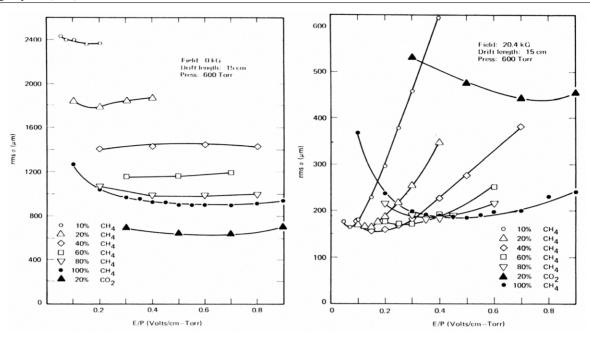


Figure 12. Transverse diffusion measured in Ar/CH₄ mixtures at 600 Torr after 15 cm drift without magnetic field B and with $B = 20.4 \,\mathrm{kG}$ [4].

For the proportional tube with wire radius a and cathode radius b, E_{op} and E_{W} are obviously radial with

$$E_{\rm op} = V/[r \ln(b/a)].$$
 (2.51)

We assume constant mobility μ for the positive ions. Therefore

$$v^{+}(t) = \mu V/[r(t)\ln(b/a)].$$
 (2.52)

For a positive ion starting at t=0 from $r=r_1$,

$$r(t) = r_1(1 + (t/t_0))^{1/2},$$
 with $t_0 = r_1^2 \ln(b/a)/(2\mu V)$. (2.53)

The maximum time for an ion to drift from a to b is

$$T_{\text{max}}^+ = (b/a)^2 t_0,$$
 as $(b/a)^2 \gg 1.$ (2.54)

The induced current I^+ is

$$I^{+} = -q E_{W} \cdot v^{+} < 0, \tag{2.55}$$

as v^+ is parallel to ${m E}$

For the integrated charge Q, one gets

$$Q^{+}(t) = \int I \, dt = \int I(1/v^{+}) \, dr = \int -q E_{w} \, dr.$$
 (2.56)

Integration from r_1 to r_2 gives

$$Q_{1\to 2}^+ = -q \ln(r_2/r_1)/\ln(b/a),$$

with
$$q > 0$$
 and $r_2 > r_1$. (2.57)

For an electron one obtains

$$Q_{1\to 2}^- = +q |\ln(r_2/r_1)| / \ln(b/a),$$

with
$$q < 0$$
 and $r_2 < r_1$, (2.58)

as v is antiparallel to E_{W} .

We shall give numbers for a typical proportional tube, assuming the following: $a=10\,\mu\text{m}$, $b=2.5\,\text{mm}$, $E_{\rm op}(r=a)=200\,\text{kV}\,\text{cm}^{-1}$, $\mu^+=1.9\,\text{atm}\,\text{cm}^2\,(\text{V}\,\text{s})^{-1}$, $v^->5\times 10^6\,\text{cm}\,\text{s}^{-1}$ and to estimate the gas amplification A, the Diethorn parametrization $\alpha=(\ln 2/\Delta V)E$ and $E_{\rm min}=V/(r_{\rm min}\,\ln(b/a))$. For an Ar/CH₄ (90/10) mixture, $\Delta V=23.6\,\text{V}$ and $E_{\rm min}=48\,\text{kV}\,\text{cm}^{-1}$ is assumed [7, p 136]. Here $r_{\rm min}$ is the starting radius for the avalanche and $E_{\rm min}$ the minimum field permitting multiplication.

We obtain $t_0 = 1.3 \text{ ns}$, $T_{\text{max}}^+ = 82 \mu \text{s}$, $r_{\text{min}} = 42 \mu \text{m}$, A = 4400.

The last electron will be collected in a very short time of about 0.6 ns, the vast majority even faster. Half of the electrons move only about $2\,\mu\text{m}$, the next 25% some $4\,\mu\text{m}$ and so on. A rough estimate of the induced electron charge signal is, therefore,

$$Q_{\text{total}}^- = q \ln(14/10) / \ln(2500/10) = 0.06q.$$
 (2.59)

Only about 6% of the total induced signal is due to the movement of the electrons, the rest comes from the ions, if one integrates over the full ion collection time, $82~\mu s$. In practice, however, one mostly uses much faster integration. The long tail in the signal caused by the very slow ion movement has to be corrected for by electronic pulse shaping to avoid pile-up at high rates. If one uses fast pulse shaping, say 20 ns integration time, only a fraction of the ion charge will be seen: an ion starting at $r_1 = a$, reaches $r_2 = 40~\mu m$ in 20 ns and induces about 25% of its charge. That means with 20 ns pulse shaping, one may expect to see an *effective charge* of about 30% of the total charge produced, of which one fifth is due to the electrons.

At higher pressure p, the same gas amplification is obtained for a lower E/p. As a consequence, ion drift is slower and therefore also the signal formation.

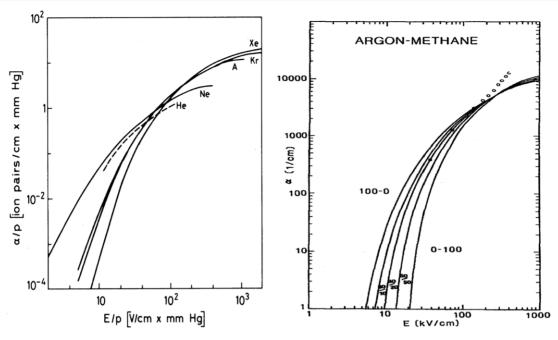


Figure 13. Measured 'first Townsend coefficient' α in noble gases [7] and calculated α in Ar/CH₄ mixtures(100-0 means 100%Ar) [41(a)], with points from measurements [41(b)]. (Left part reprinted with permission from [7]. Copyright 2008, Springer Science + Business Media. Right part reprinted with permission from [41(a)]. Copyright 1989, Elsevier.)

2.3.2. Gas amplification. In high enough electric fields, the number n of electrons will grow on a path ds by

$$dn = n\alpha \, ds, \tag{2.60}$$

where α is the *first Townsend coefficient*. As the ionization growth is proportional to the gas density ρ and the ionization cross-section depends on the instantaneous energy ε of the electrons, which is dependent only on E/ρ , α/ρ is given usually as a function of E/ρ . Some examples of measurements [40] and Monte Carlo simulation are shown in figure 13. In the region of interest to gas detectors, up to about 250 kV cm⁻¹, α rises fast with electric field. The agreement between measurements and calculation is reasonable for the lower field values. The difference at the highest fields is attributed to the fact that photo- and Penning-ionization has been neglected.

The amplification A in the detector is then

$$A = n/n_0 = \exp \int \alpha(s) ds = \exp \int \alpha(E) dE/(dE/ds),$$
(2.61)

if n_0 is the initial number of electrons, e.g. from a track segment.

In the amplification region near the wire, the field is

$$E(r) = q_s/(2\pi r \varepsilon_0), \qquad (2.62)$$

where q_s is the charge per centimetre. Therefore,

$$A = \exp \int q_s \alpha(E) \, dE/(2\pi \varepsilon_0 E^2). \tag{2.63}$$

Two approximations in particular have been used to describe practical cases.

The early Korff model [42] uses the parametrization

$$\alpha/p = A \exp(-Bp/E), \tag{2.64}$$

with empirical constants A and B depending on the gas.

In the *Diethorn approximation* [43], α is assumed to be proportional to E. One then obtains for a proportional tube with wire radius a and tube radius b

$$\ln A = (\ln 2/\ln(b/a))(V/\Delta V)\ln(V/(\ln(b/a)aE_{\min}),$$
(2.65)

where the two parameters E_{\min} and ΔV are obtained from measurements of A at various voltages and gas pressures. E_{\min} is the minimum E field to start the avalanche and $e\Delta V$ the average energy required to produce one electron. A list for E_{\min} and ΔV for various gases is given, e.g., in [7]. As E_{\min} is defined for a density ρ_0 at STP, $E_{\min}(\rho) = E_{\min}(\rho_0)(\rho/\rho_0)$. Reasonable agreement with the experimental data is obtained; discrepancies show up at high A.

The amplification $A = n/n_0$ is the mean of a Gaussian distribution with variance S^2 . If $n_0 \gg 1$ and if all n_0 avalanches develop independently, $S^2 = n_0 \sigma^2$, where σ^2 is the variance of the probability function P(j) for producing an avalanche of j electrons starting from one, if the mean is A. Distributions for single-electron amplification derived theoretically [44] and measured [45] for the strong inhomogeneous field around a thin wire approach Polya distributions, see figure 14.

For these distributions

$$(\sigma_A/\langle A \rangle)^2 = f, \quad \text{with } f \leqslant 1. \tag{2.66}$$

The limiting case f=1 is an exponential distribution (Yule–Furry law)

$$P(A) = (1/\langle A \rangle)e^{-A/\langle A \rangle}.$$
 (2.67)

Experimental results point to f = 0.6–1.0. Measurements with laser tracks [7] indicate that the rms width σ_A of a single-electron avalanche is close to the mean, as it is for the exponential distribution. Another very careful measurement also favours an exponential distribution [14, 24], except for a sharp drop at very low pulse heights.

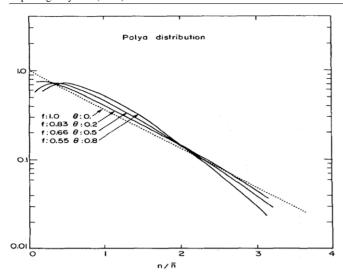


Figure 14. Polya distributions. (Reprinted with permission from [13]. Copyright 1980, Elsevier.)

For a parallel plate geometry, calculations [46] agree well with measurements [47].

Besides avalanche fluctuations, many operational and geometrical parameters influence the gas amplification, amongst which are the following:

- Geometrical imperfections. The effects will obviously depend on the geometry and the operation details. An early publication [37] gives analytic estimates of the effects of wire displacements and variations in wire diameter. In a typical geometry $dA/A \sim 2.5 dr/r$, where r is the wire radius; $dA/A \sim 9 dgap/gap$.
- Edge effects. Near edges, the electric field is reduced over distances similar to the gap between the electrode planes. It can be recovered largely by additional field shaping lines on the edges [48].
- Gas density. The Diethorn approximation gives $dA/A = -(\ln 2/\ln(b/a))(V/\Delta V)(d\rho/\rho) = -(5-8) d\rho/\rho$ typically. Pressure dependence can be corrected for, as pressure is usually well measured.
- Space charge. Due to the low velocity of the positive ions (falling off as 1/r from $> 1 \text{ mm } \mu \text{s}^{-1}$ at r = a), space charge will build up at high particle fluxes and reduce the avalanche amplification. In drift tubes, the voltage drop due to the space charge from a given particle flux is proportional to the third power of the tube radius. A smaller radius thus improves the rate capability drastically.

2.4. Calibration with laser beams

The development of TPCs and other detectors with long drift distances triggered the interest in using laser beams for calibration. They should offer several advantages: stable position in space, insensitivity to magnetic fields, no multiple scattering, narrow pulse height distribution even on short track segments without δ -electrons and easy ways to produce well defined multiple tracks.

The problem was that single photon excitation had to be excluded. Lasers in the vacuum UV are impractical and the photon energies of available lasers in the near UV, e.g. N_2 -lasers ($h\nu = 3.67 \,\text{eV}$) or quadrupled Nd–Yag lasers ($h\nu = 4.68 \,\text{eV}$), are too low to directly ionize even the lowest ionization potentials IP of complex molecules.

The first successful double photon ionization was obtained with a N_2 -laser with nickelocene (IP = 6.5 eV) as additive [49]. Soon two independent studies showed that significant ionization could be produced even in undoped gases on impurities [50,51]. A detailed interpretation was given [50], also for the fact that sometimes a linear dependence of signal versus laser intensity was observed, assuming saturation of the first step. It was demonstrated that with a special laser construction for low beam divergence sufficient ionization is obtained over long distances [51].

These first results triggered a large number of studies. Many additives were shown to be ionized with the N_2 -laser and even more with the higher photon energy of a quadrupled Nd–Yag laser ($h\nu = 4.68\,\mathrm{eV}$). Although traces of many possible candidates for two-photon ionization were detected in samples of gases from DCs, the impurities responsible for the signals in 'undoped' chambers remained unknown, except in two cases, where phenol and toluene were found to contribute [52,53]. An account of the vast amount of studies may be found in two reviews [55,56].

It turned out, as a great surprise, that adequate signals could be detected in basically all undoped chambers, with good stability over long times. It was even difficult to suppress laser ionization completely in very clean systems [54]. Therefore, operation of both TPCs and of the jet chamber at LEP was started tentatively with undoped gas mixtures. It was found that the signals remained acceptable without additives over the whole operation period of a decade, even with the lower photon energy of the N2-lasers in DELPHI. An example of reconstructed laser tracks in ALEPH is shown in figure 15. The most recent two large TPCs for STAR and ALICE have both installed very complex systems [57, 58] and installation is foreseen for the T2K TPCs. STAR is using a system of 500 beams to monitor drift velocity to 0.02% and to correct various $E \times B$ corrections and is relying strongly on them for measurements of space charge effects, especially at the higher RHIC luminosities.

The spatial stability of the laser beam can be about $100\,\mu\text{m}$; with only $100\,\text{shots}$, the mean position is thus defined to $10\,\mu\text{m}$. Besides their use in large chambers, laser beams have also found applications in numerous small-scale studies of DC performance.

2.5. Ageing of wire chambers

Since the invention of gaseous detectors ageing has been observed. It has become one of the major concerns in recent experiments, because of the enormous radiation dose detectors have to stand. Amongst the effects of *ageing* are decreasing signal, broadening energy resolution and increasing dark currents, finally electric breakdown or even broken wires. Several reviews [59] and workshop proceedings [60] give a detailed account of the immense number of studies of this subject.

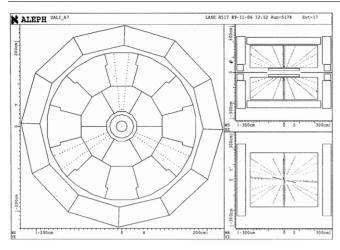


Figure 15. Reconstructed tracks from laser beams in the ALEPH TPC. (Reprinted with permission from [71(a)]. Copyright 1990, Elsevier.)

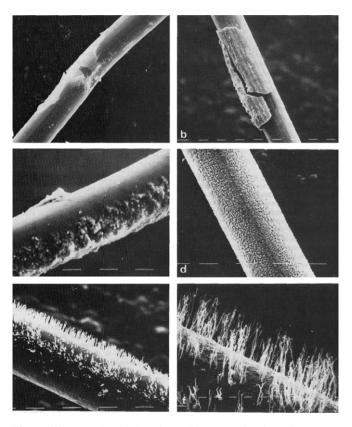


Figure 16. Examples of deposits on $20 \,\mu m$ anode wires after strong irradiation. (Reprinted with permission from [61]. Copyright 1983, Elsevier.)

As causes of the ageing effects, deposits have been found on sense wires and cathodes. Any form from smooth layers to long whiskers has been observed on the anode wires [61], see figure 16. On cathodes, thin insulating spots are usually found in discharge areas, representing signs of a Malter effect [62]: under irradiation charges build up until their field extracts electrons from the cathode through the insulator into the gas, where they initiate new avalanches. Delays of the charging-up shortening with higher irradiation rate and slow decay after irradiation is stopped support this assumption. Addition

of small amounts of alcohol or water vapour has sometimes reduced the effect, probably by providing some conductivity. Analysis of the deposits often indicated carbon compounds and also Si, and more rarely other elements, Cl, O, S.

The drop in pulse height A often follows the integrated charge load in Coulomb per centimetre wire, independent of radiation rate, but occasionally some rate dependence has been observed.

Very crudely, classical gas mixtures containing hydrocarbons have shown ageing rates of

$$\Delta A/A \sim (0.01 - 0.1)\% \, \mathrm{mC^{-1} \, cm^{-1}}$$
 for small detectors, $\Delta A/A \sim (0.1 - 1)\% \, \mathrm{mC^{-1} \, cm^{-1}}$ for large detectors.

Many qualitative explanations have been proposed for specific cases of ageing but there exists no quantitative explanation. On the other hand, the studies permit the formulation of some *general rules* for increasing the chance for a longer detector survival:

Hydrocarbons (methane, ethane, etc), very often used in the past, should be avoided for the highest radiation levels. For these, only a very restricted list of suitable gases remains: mixtures of rare gases and CO_2 and possibly N_2 , CF_4 or DME. CF_4 offers high electron drift velocities and has proven to be capable under certain conditions to avoid formation of or even to etch away deposits, in particular in the presence of minute Si impurities. But its aggressive radicals can also etch chamber components, especially glass [63]. In any case, the water content has to be kept below 0.1%, if CF_4 is used, to avoid etching even of goldplated wires. Also DME, offering low diffusion, has in some cases provided long lifetime. It was, however, proven to attack Kapton and to be very sensitive to traces of halogen pollutants at the ppb level.

- Many materials have to be avoided [59(e)], in the gas system, in the detector and during the construction: Si compounds, e.g. in bubbler oils, adhesives, vacuum grease or protection foils, PVC tubing, soft plastics in general, certain glues and many more. Details can be found in the workshop proceedings and reviews mentioned, also on suitable materials.
- *High cleanliness* has to be observed during construction, e.g. to avoid resistive spots on the cathodes. The sense wire has to be continuously checked during wiring to assure the required quality of its geometrical tolerances and of the gold plating.
- Great effort should go into the electronics to allow the *lowest gas amplification* possible.
- A *final detector module* with the final gas system components should be *extensively tested* under irradiation *before 'mass production'*. Last minute changes have often caused problems. As an accelerated test over typically <1 year is usually required for practical reasons, to obtain the full integrated charge for some 10 years of operation, an uncertainty will unfortunately remain, because a rate dependence of the ageing can not be excluded.
- TPCs are meant to see significantly lower particle rates and thus less integrated charge per centimetre of anode wire than other high-rate detectors at LHC, as already

dictated by the long drift length and resulting space charge and event reconstruction problems. Nevertheless, the beforementioned precautions are obligatory in most experiments, because radiation hardness can be deteriorated by orders of magnitude, e.g. by inadequate choice of materials or construction procedures.

3. TPC characteristics

Table 3 summarizes properties and performance of some TPCs.

3.1. Basic design—PEP4 TPC

The TPC is basically a long DC with parallel electric and magnetic fields and 3D points read-out at the end(s). The design principles shall be explained by taking the example of the PEP4 TPC, not only because this was the original TPC but also because the choices taken at that time were based on studies of many of the options, which have again been considered in later developments.

The design of the PEP4 TPC proposal is sketched in figure 17. Developed for operation at an electron-positron collider, it is in the form of a double cylinder around the beam pipe, 2 m long, centred on the interaction point and inside a solenoid. A central high voltage plane (on negative potential) supplies the field for drifting electrons towards the two read-out planes, one at each end; the drift length is 1 m. An inner and an outer field cage terminate the drift volume radially (radius $= 20 \,\mathrm{cm}$ and $100 \,\mathrm{cm}$, respectively) and define a homogeneous drift field. Each end-cap has the MWPC arranged as six sectors, each with 183 'azimuthal' anode wires. Figure 18 shows three cuts through the MWPC. The wire plane terminating the drift volume acts as a gating grid. This grid was actually only added for the second run period, because space charge problems due to background had turned out bigger than anticipated. When the gate is 'open', the electrons drift through it and also through the second grid, which acts as one of the cathodes of the MWPC. Avalanches are initiated near the anode wires (diameter = $20 \,\mu\text{m}$). The anode wires $(20 \,\mu\text{m})$ have a pitch of 4 mm and alternate with field wires (75 μ m). The field wires were considered important to reduce the crosstalk between the anode wires, as their pulse heights are used to provide PID through ionization sampling. The other cathode is formed by a printed circuit board, segmented into equidistant rows of (7.5×7.5) mm² pads below 15 of the sense wires. The number of pad rows was limited by space for the preamplifiers on the sectors and cost of the read-out. Centroid calculation of the pad pulse heights provides precise $r-\varphi$ information along the wire. The signals from all anode wires and cathode pads are sampled at 10 MHz with CCDs, which provide z-information (along the drift direction). The CCDs accept 455 analogue samples and are read out at about 20 kHz. For a track traversing the full radius, there are thus 15 points measured with 3D information and 183 dE/dx samples for PID.

A high pressure of 10 atm was specified to optimize PID with a compact design. An Ar/CH₄ (80/20) mixture was chosen, because of low electron attachment and chiefly because

of the high drift velocity u at low field E, due to the Ramsauer dip in the cross-section. These two characteristics are especially important at high pressure to avoid higher electric fields and electron attachment from oxygen impurities, which increases with the square of the pressure. The original proposal foresaw operation at p = 10 atm, $E = 0.2 \,\mathrm{V \, cm^{-1} \, Torr^{-1}}$ and 15 kG. Measurements of the transverse diffusion of single electrons as a function of E had been carried out at 600 Torr with 15 cm drift in various gases, in magnetic fields up to 20.4 kG, see figure 12. From these measurements, $\omega \tau \sim$ 10 was obtained at 600 Torr for 20.4 kG. This permitted an estimate of the $r-\varphi$ smearing due to transverse diffusion for a 1 m drift in 15 kG at 10 atm of about 1.2 mm rms for a single electron. Assuming 160 electrons, i.e. the most probable number for a 4 mm sample, contribute to the signal, the measurement error due to diffusion was estimated as $\sigma_{\rm D}$ < $100 \, \mu \text{m}$ (a very conservative estimate for the effective number of 50 would lead to $\sigma_D = 170 \,\mu\text{m}$). The specification of an average resolution of 200 μ m/point was, therefore, considered conservative and used to estimate the momentum resolution to about $\delta p/p = 0.6p\%$ for p > 1 GeV/c, well adapted to the PEP energies.

Initial space charge effects became negligible after the installation of the gating grids. Distortions due to charging-up of the field cages were strongly reduced by a reduction of the non-conducting sections and by the application of a surface coating. With the final field of $13.25\,\mathrm{kG}$ at $8.5\,\mathrm{atm}$, an average point resolution of $150\,\mu\mathrm{m}$ was achieved, the best of all large TPCs. The momentum resolution, however, remained somewhat worse than expected with about $1.0\,\mathrm{p}\%$, for reasons only partially understood [64].

It is interesting to note that for the same gas and *E*-field, at 1 atm a field of 15 kG along the drift direction reduces the single-electron diffusion $\sigma_{D,1e}$ from about 4130 to 700 μ m, a value below that for 10 atm, because of an $\omega\tau=5.8$ at 1 atm, compared with $\omega\tau=0.58$ at 10 atm. It is because of the higher effective number of contributing electrons that the higher pressure results in a smaller error in the measurement of the mean.

Excellent PID was achieved with a resolution of 3.0% for minimum ionizing particles and 2.7% for Bhabha electrons, the best value of any TPC to date [64].

3.2. Specific aspects

3.2.1. Wire chamber layout. Several parameters in the wire chamber layout may be varied to adapt to particular priorities: use of field wires, gap sizes, wire pitch and sector boundary design. Figure 18 shows the layout chosen by PEP4.

Field wires. The PEP4 TPC used field wires to reduce crosstalk between the anode wires and thus corrections to the dE/dx samples taken from them. Most TPCs followed this choice. The argument drops of course when the anode wires are no more read out, as in STAR and ALICE, because of the extreme occupancy. Dropping the field wires brings the advantage of a higher percentage of the induced charges on the pads. In the ALEPH geometry, with 4 mm gaps (anodecathode) and 4 mm anode wire pitch, pads receive about 20%

Table 3. Characteristics and performance of some TPCs.

Parameter/Experiment	PEP4	TRIUMF	TOPAZ	AlEPH	DELPHI	STAR	ALICE ^a
Operation	1982/1984	1982/1983	1987	1989	1989	2000	2009
Inner/Outer radius (m)	0.2/1.0	$\sim 0.15/0.50$	0.38/1.1	0.35/1.8	0.35/1.4	0.5/2.0	0.85/2.5
Max. driftlength $(L/2)$ (m)	1	0.34	1.1	2.2	1.34	2.1	2.5
Magnetic field (T)	0.4/1.325	0.9	1	1.5	1.23	0.25/0.5	0.5
Gas:	Ar/CH ₄	Ar/CH ₄	Ar/CH ₄	Ar/CH ₄	Ar/CH ₄	Ar/CH ₄	Ne / CO_2 / N_2
Mixture	80/20	80/20	90/10	91/9	80/20	90/10	90/ 10/ 5
Pressure (atm)	8.5	1	3.5	1	1	1	1
Drift field (kV cm ⁻¹ atm ⁻¹)	0.088	0.25	0.1	0.11	0.15	0.14	0.4
Electron drift velocity (cm μ s ⁻¹)	5	7	5.3	5	6.69	5.45	2.7
$\omega \tau$ (see section 2.2.1.3)	0.2/0.7	2	1.5	7	5	1.15/2.3	<1
Pads: Size $w \times L \text{ (mm} \times \text{mm)}$	7.5×7.5	$(5.3-6.4) \times 19$	$(9-11) \times 12$	6.2×30	\sim 7 × 7	2.85×11.5	4×7.5
						6.2×19.5	$6 \times 10/15$
Max. no. 3D points	15—straight	12	10—linear	9 + 12—circular	16—circular	13 + 32—straight	63 + 64 + 32
dE/dx: Max. no. samples/track	183	12	175	148 + 196	192	13 + 32	63 + 64 + 32
Sample size (mm atm); w or p	4×8.5 ; wires	6.35; wires	4×3.5 ; wires	4; wires	4; wires	11.5 + 19.5; pads	7.5 + 10 + 15; pads
Gas amplification	1000	50 000		3000-5000	5000	3000/1100	20 000
Gap a-p; a-c; c-gate ^b	4; 4; 8	6	4; 4; 8	4; 4; 6	4; 4; 6	2; 2; 6/4; 4; 6	2; 2; 3/3; 3; 3
Pitch a–a; cathode; gate	4; 1; 1		4; 1; 1	4; 1; 2	4; 1; 1	4; 1; 1/4; 1; 1	2.5; 2.5; 1.5
Pulse sampling (MHz/no. samples)	10/455, CCD	only 1 digitiz., ADC	10/455, CCD	11/512, FADC	14/300, FADC	9.6/400	5-10/500-1000, ADC
Gating ^c	≥1984 o.on tr.	≥1983 o.on tr.	o. on tr.	synchr. cl.wo.tr	static	o.on tr.	o.on tr.
Pads, total number	15 000	7800	8200	41 000	20 000	137 000	560 000
Performance							
$\Delta x_{\rm T}$ (μ m)-best/typ.	130-200	200/	185/230	170/200-450	180/190-280	300-600	spec:800-1100
$\Delta x_{\rm L}$ (μ m)-best/typ.	160-260	3000	335/900	500-1700	900	500-1200	spec:1100-1250
Two-track separation (mm), T/L	20		25	15	15	8 - 13/30	•
$\partial p/p^2$ (GeV/c) ⁻¹ : TPC alone; high p	0.0065		0.015	0.0012	0.005	0.006	spec:0.005
dE/dx (%) Single tracks/ in jets	2.7/4.0		4.4 /	4.4 /	5.7/7.4	7.4/7.6	spec:4.9/6.8
Comments		a in single PCs	chevron pads	circular pad rows	circular pad rows	No field wires	No field wires
		strong $E \times B$ effect	•	•	•	>3000 tracks	≤ 20000 tracks

Expected performance.
 a = anode, p = pads, c = cathode grid.
 o. on tr.: gate opens on trigger; cl.wo.tr. : opens before collision and closes without trigger; static : closed for ions only (see text).

Table 3. Continued.

Parameter/Experiment cont.	NA35	EOS/HISS	NA49 VTX	NA49 MAIN	CERES/NA45	HARP	T2K ^a
Operation	1990	1992	1995	1995	1999	2001	2009/10
Inner/Outer radius or L/W (m)	2.4/1.25 (L/W)	1.5/0.96 (L/W)	$2.5/1.5$ (L/W); $2 \times$	$4/4 (L/W); 2 \times$	0.6/1.3; L = 2	0.1/0.41	$2.2/0.7$ (H/L); $3 \times$
Max. driftlength $(L/2)$ (m)	1.12 vert.	0.75 (H)	0.67 vert.	1.1 vert.	0.7 rad.	1.6	0.9 W
Magnetic field (T)	0	1.3	1.5	0	$B_z < 0.7; B_r < 0.3$	0.7	0.2
Gas:	Ar/CH ₄	Ar/CH ₄	Ne/CO ₂	Ar/CH ₄ /CO ₂	Ne/CO ₂	Ar/CH ₄	Ar/CF4/i-C4H10
Mixture	91/9	90/10	90/10	90/ 5/5	80/ 20	91/9	95/3/2
Pressure (atm)	1	1	1	1	1	1	1
Drift field (kV cm ⁻¹ atm ⁻¹)	0.12	0.12	0.19	0.175	0.2-0.6	0.111	0.2
Electron drift velocity (cm μ s ⁻¹)	5	5.5	1.3	2.3	0.7-2.4	5.2	7
$\omega \tau$ (see section 2.2.1.3)	0	0.5	1	0		3.3	0.7
Pads: size $(w \times L, mm \times mm)$	5.5×40	8×12	$3.5 \times (16, 28)$	$(3.6, 5.5) \times 40$	10 chevron	6.5×15	6.9×9.7
Max. no. 3D points	60 + 30	128	<150	90		20	72×3
dE/dx: Max. no. samples/track	60	128	<150	90		20	72×3
Sample size (mm atm); w or p	40; pads	12	16, 28	40		15	9.7
Gas amplification	•	3000	20 000	5000	8000	20 000	$\sim \! 1000$
Gap a–p; a–c; c–gate ^b		4; 4; 6	3,2;	2,3; 3;6	3;3;6	5;5;6	0.128
Pitch a–a; cathode; gate	4; 1; 2	4; 1; 2	4; 1; 1	4; 1;1	6; 2; 2	4; 2; 2 stagg.	
Pulse sampling (MHz/no. samples)	12.5 /	10/256, SCA	/512	/ 512		10/>300, FADC	/512 SCA
Gating ^c		o. on tr.	o. on tr.	o. on tr.	o. on tr.	o.on tr.	none
Pads, total number	11 000	15 000	74 000	108 000	78 000	4000	125 000
Performance							
$\Delta x_{\rm T}$ (μ m)-best/typ.	300-800	300	150	150	230/340	600-2400	600 (1m drift)
$\Delta x_{\rm L}$ (μ m)-best/typ.	250-450				dr = 400/640	3.5	,
Two-track separation (mm)	18	25		10	,		
$\partial p/p^2$ (GeV/c) $^{-1}$: TPC alone; high p		1			1	0.2/0.45-0.50	spec: <10;
dE/dx (%): single tracks/in jets	/6	/ 4	<4 : VTX + Main			16	spec: <10/
Comments	B = 0 only pad r.o.	only pad r.o.	Kr ^m calibration only pad r.o.	up to 1200 tr. only pad r.o.	Radial TPC No field wires	el. crosstalk	Micromegas r.o.

a Expected performance.
b a = anode, p = pads, c = cathode grid.
c o. on tr.: gate opens on trigger; cl.wo.tr. : opens before collision and closes without trigger; static : closed for ions only (see text).

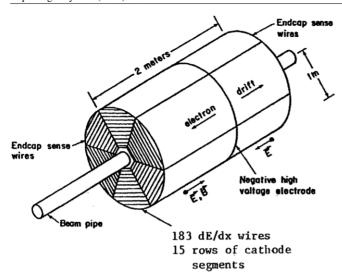


Figure 17. Sketch of the PEP4 TPC. (Reprinted with permission from [PEP4(d)]. Copyright 1983, IEEE.)

of the anode charge with the field wires (which get about 40%) and would receive about 35% without them [65]. In the inner sectors of STAR even >40% is received by the pads, because of the small anode–cathode gap of 2 mm.

Dropping the field wires avoids problems from corona discharges on the frames, where anode and field wires are soldered. This was actually considered the major argument against adding field wires.

On the other hand, operation without field wires demands a higher voltage on the anode wires for the same gas amplification.

Pads. The PEP4 TPC and most later TPCs have straight pad rows. As track angles not normal to the pad row cause an additional error for the spatial resolution, ALEPH and DELPHI have chosen circular pad rows in the cylindrical geometry of a collider. The choice of the size of the pads depends on many parameters including cost. The width along the pad row, i.e. along $r-\varphi$, is usually smaller than the length. ALEPH decided for longer pads to improve resolution for the highest momenta, DELPHI for short pads to improve performance for tracks from secondary vertices. The effect of the length on spatial resolution is complex, see below. For best space resolution, the ratio 'pad width/gap (anode-cathode)' should be about 1–2 [120, 121]. For larger ratios, precision is lost for tracks passing above the centre of a pad. For lower ratios, the precision is limited by the lower signal-to-noise ratio. To reduce occupancy, STAR and ALICE chose thinner pads (2.85 mm and 4 mm, respectively) for the inner circle of sectors and NA49 for the Veretx TPC (3.5 mm).

Gaps and wire pitch. The gap (anode-pad) has been chosen as 4 mm for most TPCs, but smaller for some: 3 mm for CERES, NA49 and ALICE outer chambers, even only 2 mm for the inner chambers of STAR and ALICE, matching their smaller pad width.

A pitch of the anode wires of 4 mm has been used in almost all TPCs, as it provides operational stability and a reasonable width for the dE/dx samples. An exception is made again by

the inner ALICE chambers with a pitch of 2.5 mm (without field wires), matching the thin gaps.

The pitch of the cathode and gating grids is 1 or 2 mm. The 1 mm grids show smaller $E \times B$ effects on the spatial resolution but exert larger forces on the frames. For the gating grid, a 1 mm pitch brings the obvious advantage of lower gating voltage.

3.2.2. Sector boundaries. The sector boundaries formed by the frames supporting the wire planes represent dead zones in the endcaps. A higher number of sectors per 2π results in more dead zones but smaller track angle errors for the $r-\varphi$ measurement. Larger TPC radii demand more sectors per 2π to remain with stable wire length. One has to choose between full insensitivity for certain φ -segments, using the same radial boundaries over the full radius (e.g. PEP4, DELPHI (6 sectors/ 2π), STAR(12), ALICE(18) or reduced sensitivity over larger φ -segments, by either using boundaries with steps along $r-\varphi$ (already discussed for PEP4 but not implemented) or by rotating in φ a second outer circle of sectors with respect to the inner one (e.g. ALEPH).

In addition to complete loss of information along the boundaries there is a serious loss inside the sensitive zone due to a reduction of gas amplification over more than a centimetre near the boundaries. Figure 19 shows an early measurement of this effect and also the corrective measure [48]. Additional guard strips on the wire support frames on a suitable potential can recover some 8 mm with full gain.

Because of their long drift length, TPCs *3.2.3. Gating.* are particularly prone to space charge effects. For the PEP4 TPC it was estimated that a charge deposition of 10 GeV s⁻¹ equivalent to 5000 m tracks/s, would cause detectable field distortions. Background turning out higher than expected, a gating grid had to be added. Most later TPCs used a similar geometry, see figure 20. A grid with wires stretched parallel to the anode wires with a pitch of 1-2 mm is mounted 4-8 mm above the shielding grid. To close the gate, a differential potential $V = V_G \pm \Delta V$ is applied to alternating wires. For the ALEPH specifications (Ar/CH₄ at 1 atm) the following results were obtained [66]. A gating grid with 1 (2) mm wire pitch is fully transparent for a V_G , which produces a field between the gating grid and the cathode about twice the value of the main drift field. The gate is closed for the ions with $\Delta V = \pm 20$ (40) V, both without magnetic field B and with B = 1.5 T. For electrons, it is also closed for B = 0, but it opens when the field is raised. For B = 1.5 T, $\Delta V = \pm 190 \text{ V}$ is required for a grid with 2 mm pitch. At $\Delta V = \pm 40 \text{ V}$, the transparency for electrons is about 75%. The big difference in the behaviour of ions and electrons in a strong magnetic field stems from the fact that in the gas mixture used $\omega \tau \gg 1$ for electrons and $\omega \tau \ll 1$ for ions. Electrons, therefore, follow closely the magnetic field lines, whilst ions follow mainly the electric field lines.

The gating behaviour was measured without magnetic field also at 8.5 atm in Ar/CH₄ [67]. A gate with 1 mm pitch showed full transparency for a field between cathode and gating grid equal to 1.9 times the drift field. The gate was closed for $\Delta V = \pm 100 \, \text{V}$.

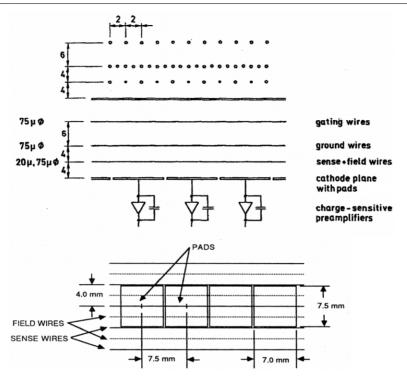


Figure 18. Schematic wire and pad layout in the PEP4 TPC proportional chambers [PEP4(e)].

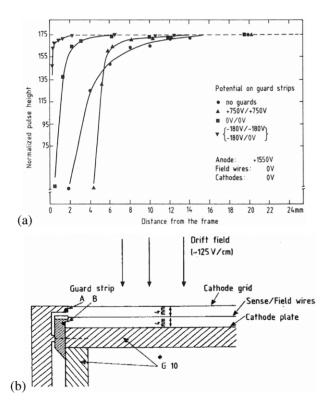


Figure 19. DELPHI TPC–sector boundary: (*a*) pulse height variation near the boundary with and without guard strip. (*b*) Details of the frames and guard strips at the boundary. (Reprinted with permission from [48]. Copyright 1985, Elsevier.)

The gate may be used in different ways. The simplest is the *static mode*, which blocks all ions but lets most of the electrons pass, with a signal loss of about 25%. DELPHI found this acceptable for operation at LEP. In the second mode, *opening*

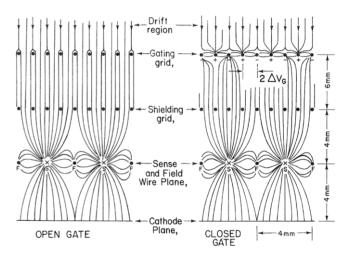


Figure 20. Schematics of the field lines for an open and closed gating grid. (Reprinted with permission from [5]. Copyright 1982, World Scientific.)

on a trigger, the gate is normally closed but opens for the full drift time for a triggered event. This mode is used in the vast majority of TPCs and is foreseen for ALICE. Depending on the speed of the trigger decision and switching, the information of a small part of the drift volume is lost; in the case of the STAR trigger, opening takes $2.1 \,\mu\text{s}$, resulting in $12 \,\text{cm}$ loss of the $2.1 \,\text{m}$ drift length. This loss is avoided in the *synchronous mode*, in which the gate is opened shortly before each bunch crossing and closed after a negative trigger decision or left open for the electron drift time after a positive trigger. ALEPH applied this mode at LEP, where the spacing of bunch crossings was $22 \,\mu\text{s}$.

The TRIUMF TPC used a different gate arrangement. Each of the anode wires is mounted with its pad row inside

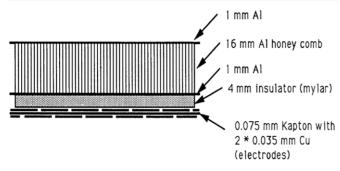


Figure 21. Schematic cut through the field cage of the ALEPH TPC. (Reprinted with permission from [5]. Copyright 1982, World Scientific.)

a rectangular cathode, sampling the track ionization through a 6 mm slot. Above each slot are mounted two grids (wire pitch 0.6 mm) 1.6 mm apart. The gate is normally closed, and opened within 200 ns on a trigger. The opening is achieved by applying -48 V on the outside grid and +3 V on the grid nearer to the anode. The dual grid has a transparency of 90% and deteriorates the spatial resolution by about 15%.

3.2.4. Field cages. The field cages have to provide a very homogeneous electric field parallel to the magnetic field axis. Their construction is complicated by conflicting requirements: high mechanical and electrical precision and stability of geometry and alignment with the cathode and the MWPC on the one hand, minimized material and thickness on the other hand. Depending on the drift length and gas choice, voltages up to 100 kV are required. The basic principle is a linear voltage degrader composed of closely spaced parallel rings connected to a chain of high precision resistors. The cathode is connected to negative high voltage and the MWPC kept near ground potential.

For the insulation towards the outside two techniques are employed, using either a solid or a gaseous insulator.

The solid insulator can be much thinner geometrically, allowing smaller detectors beyond the TPC. It usually results, however, in more material in terms of radiation length. One example of this type is the choice for ALEPH field cage, 4.4 m long and 3.8 m in diameter, see figure 21. It has a double layer of electrodes (35 μ m Cu) separated by a Kapton foil (75 μ m) on a 4 mm thick insulator, composed of 120 layers of Mylar wound helically and glued together. An aluminium honeycomb 16 mm thick and sandwiched between two 1 mm aluminium foils provides the self-supporting structure. The total thickness is 4.8% of a radiation length.

An example of a field cage with gaseous insulator is the one of ALICE. The choice of a low mass gas mixture with Ne demands a very high voltage of $100\,\mathrm{kV}$ for the $2.5\,\mathrm{m}$ drift length to obtain an acceptable drift velocity, still only about half that of the more common Ar/CH₄ mixtures. The electric field is formed by 166 aluminized Mylar strips wound around 18 support rods connecting the central cathode plane with an end plate. One rod contains the chain of resistors $(7.5\,\mathrm{M}\Omega)$ and clamps to fix the strips. The rods lie in line with the dead zones between the read-out chambers. They are clamped to the wall of a vessel containing CO_2 as gas insulator, $15\,\mathrm{cm}$ thick.

Both walls of this vessel are built with Nomex honeycomb sandwiches with prepreg and Tedlar skins. The wall towards the drift volume has Al strips on both sides, connected to a separate voltage degrader chain and the other wall has a continuous $50~\mu m$ Al foil on both sides and provides the ground termination.

In all designs, insulating surfaces have to be minimized, especially between the conducting strips, to avoid charging-up effects and resulting field distortions. Such a problem on the initial PEP4 field cage was cured by covering it with a thin layer of polyurethane, thus providing some small conductivity. Excellent alignment with the central electrode and the read-out chambers is obviously important.

Central cathode. The central cathode for TPCs at colliders has to be thin, as it lies in the symmetry plane around the interaction point. It should be flat and not deform with time. Several designs have been used: stretched stainless steel mesh (TOPAZ), carbon-coated Mylar foil (ALEPH), 25 μ m aluminized Mylar (ALICE), 70 μ m carbon-loaded Kapton (STAR) and aluminium honeycomb sandwich (DELPHI). In STAR, the membrane is secured under tension to an outer support hoop, which is mounted on the outer field cage. The membrane has just a central hole to pass the inner field cage but without any mechanical coupling to it except for a single electrical connection. Narrow aluminium strips are attached to each side of the membrane to provide a target with low workfunction for the UV laser.

3.2.5. Calibration. In addition to the usual calibration procedures of wire chambers during and after construction, regular calibration and monitoring during operation is required for optimum performance of a TPC. Various techniques have been used.

The electronics is calibrated by pulsing the grid wires (e.g. PEP4, TOPAZ and STAR) or the field wires (e.g. ALEPH and DELPHI) and recording the response of the wires and pads at various pulse heights to cover the dynamic range. This calibration is important for the $r-\varphi$ and dE/dx measurements. It also provides relative timing inside a sector.

Radioactive sources in the endplanes were used in several TPCs (PEP4, TOPAZ, DELPHI): rods embedded in the endplane can be moved to provide a point source through small holes to each wire in the 'ON' position. Three such rods per sector define three absolute gain calibrations of the wire and its electronics as well as the slope and curvature of the gain along the wires.

Actual event tracks are used for relative sector to sector timing; the time of tracks passing through the amplification zone defines the zero time for the drift time measurement. Time association of hits on wires and pads gives the relative time offset between them.

Lasers have consistently been used to measure the drift velocity to a precision down to 2×10^{-4} . Laser systems with from three beams per sector to a total of almost 500 beams in STAR have been implemented. They have also been of great help in the study of all kinds of distortions in the tracking due to space charges, insulator charging up on the field cages and misalignments.

3.2.6. Choice of gas mixture. There exists no universally 'ideal gas' for a TPC. The physics to be studied defines the layout and a number of conflicting requirements for the gas. The gas mixture has to be optimized for each particular case.

The PEP4 collaboration opted for a high pressure TPC to optimize dE/dx resolution in a compact detector. As mentioned before, an Ar/CH₄ mixture offered two advantages: a high drift velocity of electrons at low electric field and low electron attachment to oxygen impurity, which is proportional to the square of the pressure. The 80/20 mixture was preferred over the 90/10 and 95/5 mixtures because of a longer 'plateau' of stable operation. Prototype measurements had shown a length of 600 V, 400 V and 230 V, respectively, for the three mixtures.

The two advantages mentioned and perhaps the fact that Ar/CH₄ had become well understood lead to the use of P20 (80/20 mixture) and P10 (90/10) in most later TPCs. P20 offers higher electron velocity but P10 a maximum of the drift velocity at lower electric field. Operation on the maximum is advantageous for stability reasons.

ALICE opted for Ne/CO₂/N₂ (90/10/5). Neon was chosen for its low radiation length and hence low multiple scattering. CO_2 replaces CH_4 as quencher to avoid common wire chamber ageing, although it is not clear that the risk is high for the anticipated radiation loads. As CO_2 is a weaker quencher than CH_4 , N_2 is added to increase the stable gain limit. This ALICE mixture, however, requires an electric field several times higher than that required for P10 or P20 to achieve a similar drift velocity. As on the working point there is a steep dependence of the drift velocity on temperature, a very tight limit of $\pm 0.1^{\circ}$ has been specified for the temperature control. Due to the lower Z, charged particles produce in Ne a factor of 2 less ionization than in Ar. The gas amplification has to be raised to about 20 000 to achieve the same signal/noise ratio.

For the Linear Collider (LC) TPC projects various mixtures are under consideration. The toughest demand for the TPC is the point resolution <140 μ m for drift lengths up to 2.5 m. The choice of the gas may depend on the read-out technique chosen but it is clear that low transverse diffusion in a high magnetic field of about 4T will be an essential requirement. One of the candidates, the so-called T2K mixture (Ar/CF₄/i-butane 95/3/2), offers a high drift velocity for electrons of 73 μ m ns⁻¹ at a moderate 200 V cm⁻¹ electric field, a relatively low longitudinal diffusion constant of 250 μ m cm^{-1/2} and a large $\omega \tau \sim 20$ at 5T, reducing the transverse diffusion coefficient to \sim 20 μ m cm^{-1/2}. From the simulation results [25] one would in fact expect a rather similar performance from the 'standard' mixture Ar/CH₄(80/20).

3.2.7. Read-out electronics. The development of TPCs is intricately linked to developments in electronics. In particular, the implementation of typically several hundred analogue samples on each channel requires special designs. The first TPC (PEP4) was made possible by the appearance on the market of CCDs, which could be used as linear analogue shift registers, with adequate performance and at acceptable cost for a system with about 16 000 read-out channels. We shall describe this system and the one from ALICE, as a recent design for about 500 000 channels.

PEP4 electronics. The PEP4 TPC demanded charge and timing information from both wires (for dE/dx) and cathode pads (for centroid finding) [68]. To reduce input capacitance and thereby noise, the preamplifiers are mounted on the backside of the TPC endplates inside the gas volume. Their signals are sent via 100 feet 50 Ω coax cables to remote amplifier/shapers each of which drives a CCD. The CCD input is sampled and pushed through the 445 buckets at 10 MHz, memorizing a 44.5 μ s history of analogue drift information. During read-out, the clock is changed to 20 kHz and the outputs of the 16 000 channels are digitized (to 9 bits) in parallel. A threshold is set at about four times rms noise and only non-zero data are transferred to buffer memories and read to tape.

The preamplifiers are charge sensitive FET integrators of low noise (<600 electrons rms on the pads and <1000 on the wires), low power dissipation (<120 mW) and occupying only 1.2 cm², an achievement at that time. Pulse shaping was chosen as long as possible to improve the signal and reduce the noise (mainly from series noise in the preamplifier). On the other hand, for good reconstruction efficiency in jets, clean separation of tracks within $1 \mu s$ was demanded. This led to the choice of a peaking time of 250 ns. The shaper (1 RC differentiator and 5 RC integrators with the same time constant) delivered a response to a delta impulse close to a Gaussian. It also provided pole-zero cancellation of the 5 μ s decay time of the preamplifier and partial correction of the 'ion tail', with a RC differentiator (0.4 μ s), keeping the amplitude below 4% of its peak value after 600 ns. The pulse is broadened by diffusion and track inclination. Because of the slow signal formation, the measured signal is only about one third of the total charge at 10 atm (about 60% at 1 atm) with 250 ns shaping time.

A sampling frequency of 10 MHz was chosen. This guarantees a good determination of the signal charge from the pulse area and of the time centroid calculated from some five samples, with small errors due to the asynchronous sampling phase, the trigger for the start of the sampling sequence being provided by other detectors. The threshold set after the digitizer corresponds to about 3–5% of the signal peak and its effect can be corrected for. With the electron drift velocity of 5 cm μ s⁻¹ finally used and a maximum drift length of 1 m, an event was fully contained in some 200 CCD buckets.

A powerful testing scheme was implemented to calibrate each of the stages from the preamplifier to the digitizers. Electronic gain could be monitored to better than 1%.

For a gas amplification of ≤ 2000 , the electronic noise was about 1% of the signal for a minimum ionizing particle, resulting in a contribution to the spatial resolution from the pads of $60 \, \mu \mathrm{m}$.

The group was proud to keep the mass construction cost to 114 \$/channel, the cost including overhead to 280 \$/channel.

ALICE electronics. For the construction of this system with about 560 000 channels and the detection of events with extremely high multiplicity, novel developments were required [69]. To handle high track densities, special attention has been given to excellent ion-tail cancellation and baseline subtraction.

Only pad signals are read out. They are transmitted via Kapton cables to front-end cards (FECs) located some

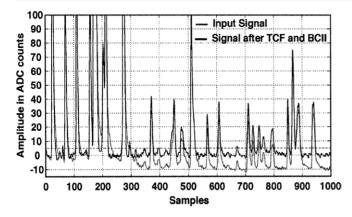


Figure 22. Performance of the ALTRO chip of the ALICE TPC: pulses from a high-multiplicity cosmic ray event before and after tail cancellation and baseline restoration. (Reprinted with permission from [69(b)]. Copyright 2004, Elsevier.)

10 cm away in a structure mechanically decoupled from the TPC endplate. Each channel in the FEC consists of two parts: (a) an analogue ASIC (PASA) with an amplifier and semi-Gaussian shaper and (b) a mixed-signal ASIC (ALTRO), including a 10-bit 25 MSPS ADC, baseline subtraction, a tail cancellation filter (TCF), zero suppression and a multi-event buffer. The shaping time is 190 ns FWHM, the sampling rate of the ADC 5–12 MHz. The power consumption of the whole FEC, including a board, controller is <100 mW per channel. This is less than what required by the PEP4 preamplifier alone and is one clear indication for the extraordinary progress of the electronics in the last 30 years.

The first component of the ALTRO chip is the baseline correction-1 to perform channel-to-channel gain equalization and to correct non-linearity and low-frequency baseline The next processor is an 18bit, fixed-point arithmetic, third-order TCF, designed to suppress the ion tail to less then 1 LSB from the baseline, within 1 μ s after the signal With fully programmable coefficients, independent for each channel, the circuit can cancel a wide range of tail shapes and correct channel-channel fluctuations. The next block, the baseline correction-2 (BC 2), applies a correction based on a moving average of samples within a window and removes non-systematic perturbations of the baseline that are superimposed on the signal. At the output a zero suppression unit discards all data below a programmable threshold, except for a programmable number of pre- and post-samples around each pulse. The output is sent to a memory with storage capacity of up to eight events of 1000 words.

Isolated cosmic ray events in a prototype were used to tune the parameters, employing the TCF to cancel the tail up to 3 μ s and the BC 2 to remove the long undershoot and a secondary spike after 25 μ s due to the acceleration of the positive ions near the grid wires. Figure 22 shows the impressive result obtained for a high-multiplicity cosmic ray event, showing the pulses before and after correction through the TCF and the BC 2.

Other specifications for the electronics are dynamic range = 900:1; Crosstalk <1%; maximum read-out rate: 200 Hz for Pb–Pb collisions with up to 700 Mbytes/event and 1 kHz for p–p collisions. An occupancy of up to 50% near the inner TPC radius should be handled.

The radiation load in the TPC is supposed to be relatively low: 1 krad + 10¹¹ neutrons cm⁻² over 10 years. Standard radiation-soft technologies have, therefore, been used. Special attention has, however, been paid to protection against damage from single event upsets by applying redundancy and solutions to detect and cure possible upsets in the configuration of the programmable devices.

3.3. Performance

3.3.1. Spatial resolution. The avalanches on the anode wires induce pulses on the cathode pads below and the $r\varphi$ coordinate is obtained from the centre-of-gravity determination of the pulse heights from adjacent pads.

Pad response. The charge distribution induced by a pointlike avalanche on an anode wire is called the intrinsic pad response function (prf). It is determined by the geometry of the proportional chamber, especially the gap g (anode–cathode). When measured with pads, the prf also depends on the pad size [120]. In addition, there is a dependence on the actual avalanche geometry—how much the avalanche surrounds the wire—which determines the path of the positive ions to the various electrodes. This is normally neglected but simulations clearly show the effect [70] and also some measurements with the ALTRO chip in ALICE [69(b)].

As an example, numbers for the PEP4 geometry are given: with an anode wire pitch of 4 mm, a gap (anode–cathode) of 4 mm and a pad size of $(7.5 \times 7.5) \, \text{mm}^2$, the measured width of the intrinsic prf is $\sigma_0' = 3.6 \, \text{mm}$.

For the many avalanches resulting from the track ionization, there are several effects broadening the distribution. The collaborations have used different parametrizations; we shall follow the one by ALEPH. The actual prf is described by

$$\sigma_{\rm prf}^2 = \sigma_0^{'2} + \sigma_{\rm D}^{'2} z + \sigma_{\alpha}^{'2} ({\rm tg}\alpha - {\rm tg}\psi)^2 + \sigma_{\beta}^{'2} {\rm tg}^2 \beta, \qquad (3.1)$$

where α and β are the angles between the track and the normal to the wires and the pad row, respectively, and ψ is the angle describing the $E \times B$ effect in a magnetic field (see figure 23); z is the drift distance.

Although in most TPCs straight pad rows follow the wire orientation, we consider the general case, which includes the circular pad rows of ALEPH and DELPHI. This has the advantage to clearly separate different effects. The errors are added quadratically, as they are assumed independent.

- The first contribution contains the width of the intrinsic prf, in addition the error from noise and calibration.
- The second term describes the broadening due to diffusion.
- $E \times B$ effect. The third term has two origins. Due to the track angle α , the avalanches are spread over a wire segment d tg α , if d is the pitch of the anode wires. In addition, there is a spread caused by $E \times B$: near the gating and cathode grids and near the anode wire, the electrons do not drift parallel to the magnetic field [73]. The deviations are described by ψ such that d tg ψ corresponds to the actual smearing along the wire due to the sum of all three deviations, see figure 24. It is this whole third term, which

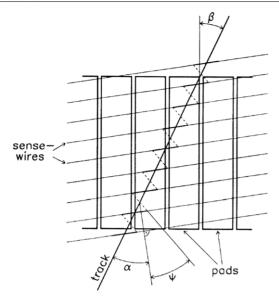


Figure 23. Definition of the track angles and the Lorentz angle ψ from $E \times B$. (Reprinted with permission from [71(b)]. Copyright 1991, Elsevier.)

one tries to suppress using read-out techniques without anode wires, see section 5. The effect could be reduced somewhat in a standard TPC by a smaller pitch of the anode wires and the wires of the cathode (which might even be replaced by a mesh) and of the gating grid. But there are limits set by electrostatic stability and mechanical forces. A strong $E \times B$ effect was observed for the first time in the TRIUMF TPC, because in the PEP4 TPC the effect was largely suppressed by the high pressure and the lower magnetic field during the early runs.

 The last term stems from the fact that several anode wires are contributing to the pad signals, each one giving a different centre of gravity. The smearing is proportional to the pad length. The error due to this term for stiff tracks is significantly reduced with circular pad rows.

The pad width w is usually chosen such that for the smallest prf width $= \sigma'_0$, two to three pads will have signals well above noise, which corresponds to a ratio w/g of about 1–2. In this case, the signal/noise ratio is near optimum [120] and it turns out that the prf is well approximated by a Gaussian with a σ close to the gap value g:

$$P_i(x) = C \exp(-(x - x_i)^2 / 2\sigma^2),$$
 (3.2)

where x is the position of an avalanche and x_i and P_i are the centre and pulseheight of the ith pad, respectively [121, 122].

From three-pad clusters P_1 , P_2 and P_3 , the prf width is deduced:

$$(w/\sigma)^2 = \ln(P_2^2/P_1P_3). \tag{3.3}$$

Measurements of clean high momentum tracks as a function of drift length and track angles α_w and α_p permit separation of the various terms and confirm the functional dependence. The Lorentz angle was measured as $\psi=32^\circ$ in ALEPH with 1.5 T.

Resolution in $r\varphi$. For the azimuthal coordinate resolution $\sigma_{r\varphi}$ one expects qualitatively the same dependence on drift distance and track angles as for the prf, although statistical effects have to be considered. In fact, a corresponding parametrization has been used by ALEPH [71].

$$\sigma_{r\omega}^2 = \sigma_0^2 + \sigma_D^2 z + \sigma_\alpha^2 (tg\alpha - tg\psi)^2 \cos^2 \alpha + \sigma_\beta^2 tg^2 \beta. \quad (3.4)$$

For stiff tracks, the angle β is close to zero, α varies from -30° to +30°. In a test setup, $\sigma_0 \leq 80 \,\mu \text{m}$ was measured, including electronic noise, digitization and errors in the calibration. ALEPH measured for the transverse diffusion term: σ_D = $35 \,\mu\text{m}\sqrt{z}$ at B = 0 and $5 \,\mu\text{m}\sqrt{z}$ at $B = 1.5 \,\text{T}$, with drift length z in cm. This agrees with expectations for 3 cm long pads, about 90 electrons per centimetre along the track and a transverse diffusion for single electrons of 600 μ m \sqrt{z} . For the α -term, $\sigma_{\alpha} = d/\sqrt{12N_{\rm eff}} = (1.15 \, {\rm mm})^2/N_{\rm eff}$ is expected, where d is the anode wire pitch and N_{eff} is an effective number of electrons statistically contributing, taking into account the The fact that for high momentum effects of clustering. tracks no significant dependence of $\sigma_{r\varphi}$ on α was observed is explained with declustering by diffusion for the long drift distances and thus increased $N_{\rm eff}$. This effect had been observed earlier [20], see section 2.1.3. The angle dependence of the β -term as measured with tracks from hadronic Z^0 decays is shown in figure 24. It gives $\sigma_{\beta} = 2.3$ mm and is the dominant contribution above a few degrees. For isolated tracks from leptonic Z^0 decays an overall resolution of $\sigma_{r\varphi} = 173 \,\mu\text{m}$ has been obtained.

For the DELPHI TPC, the Lorentz angle is $\psi=34^\circ$ [72(b)]. The resulting dependence of the resolution on $\alpha_{\rm w}=\alpha$ is presented in figure 25. The minimum at $\alpha=-34^\circ$ is just off-scale. The effect of $\beta=\alpha_{\rm p}$ is also shown, up to larger angles than in figure 24.

Longitudinal spatial resolution. The longitudinal (axial) position z of track points is calculated from a fit to a few time buckets. With increasing dip angle, the distribution of the arrival times of the electrons is stretched significantly. This reduces the longitudinal resolution, as can be seen from the measurements by ALEPH [71], figure 24.

3.3.2. Particle identification—dE/dx The principal idea of the TPC is the combination of tracking and PID in a single detector. As described in section 2.1.2, gas ionization by a particle depends only on its velocity and the square of the charge and not on its mass. PID can, therefore, be achieved by simultaneously determining the particle momentum and its velocity from the ionization. In the present context, when using the term dE/dx, because this is commonly done, we always mean the localized ionization deposited near the track.

To understand the possibilities and problems of the dE/dx measurement, we look at the results from the PEP4 TPC [15, p 298], figure 26. For particle separation, precise knowledge of two parts is required: the general dependence of dE/dx on the velocity and the ionization deposit along a specific track. The ionization loss drops as $1/\beta^2$ for low β , goes through a minimum for $\beta\gamma = 3-4$, then rises logarithmically in $\beta\gamma$ ('relativistic-rise' region) until it reaches a plateau. PID

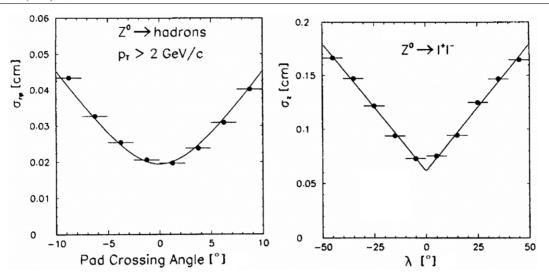


Figure 24. Spatial resolution measured with the ALEPH TPC: $r\varphi$ -resolution as a function of pad crossing angle in $r\varphi$ and z-resolution as a function of dip angle λ . (Reprinted with permission from [71(b)]. Copyright 1991, Elsevier.)

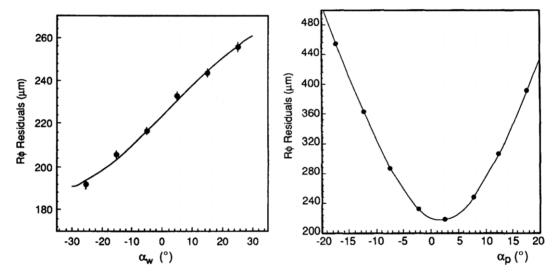


Figure 25. Spatial resolution measured in the DELPHI TPC, as a function of wire and pad crossing crossing angles, α_w and α_p . (Reprinted with permission from [72(b)]. Copyright 1992, Elsevier.)

is easy in the $1/\beta^2$ part but difficult in the relativistic-rise region. The dE/dx curve and in particular the slope of the relativistic-rise region and the plateau depend mainly on the gas and its pressure but also on the method used to measure the ionization. This function has to be determined from fitting the dE/dx measurements for well-identified particles in different $\beta\gamma$ regions, such as Bhabha electrons, cosmic muons and minimum ionizing pions.

To measure the ionization loss of a particular track with reasonable precision, many samples have to be taken on the track. This problem was first considered in [74]. A detailed study based on the photoabsorption ionization (PAI) model [75] concluded that the ionization resolution was continuously improving with increasing number of samples for a given track length. This conclusion was obtained from a likelihood treatment of the samples for different masses. Although this method is considered to be the most precise in principle, most groups have used the simpler so-called 'truncated mean' method. Figure 27 shows on the left typical distributions [76],

also called *straggling functions*, of the ionization loss in a sample of 7.5 mm in the ALICE gas: Ne/CO₂/N₂ (90/10/5). The mean and the variance are ill-defined. A more reasonable description is given by the most probable ionization $I_{\rm mp}$ and the FWHM. The procedure most widely used is the calculation of a *truncated mean*, which we denote as I, as the mean of the lowest p% of the pulse heights. The cut reduces the effect of fluctuations due to the long tail. Values of p=40–80 have been used. The spread of I, $\sigma(I)$, is not very sensitive to p in this range. Figure 27 shows on the right the distributions for p=65%, well described by a Gaussian.

Excellent agreement between experiment and theory is achieved, both for the straggling functions and the truncated means, but only after inclusion of diffusion (causing some correlation between adjacent samples) and amplification fluctuations [65]. Their effect on the straggling function is seen in figure 28. When treating different sample lengths x, because of different pad lengths or track angles, the naïve assumption of proportionality with x is incorrect, see figure 29, which shows

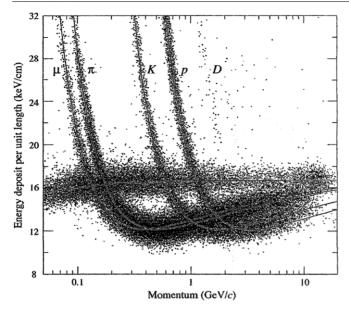


Figure 26. PID from ionization measurements (dE/dx) in the PEP4 TPC. (Reprinted with permission from [15]. Copyright 2008, The Regents of the University of California.)

calculated straggling functions for different sample lengths x in argon. As these distributions are normalized to x, the mean loss $\langle \Delta/x \rangle$ is the same, but the most probable loss changes. This means that straggling functions cannot be scaled with a single parameter (e.g. $I_{\rm mp}$). It turns out, however, that a 2-parameter scaling is adequate [77].

The quality of PID is determined by the *separation power* D with

$$D = (I_{A} - I_{B})/[\sigma(I_{A}) + \sigma(I_{B})]/2,$$
 (3.5)

i.e. the separation of particle types A and B expressed in number of standard deviations. For the PEP4 TPC, the best K- π separation obtained was about $3\sigma(I_\pi)$ with $\sigma(I_\pi)/I_\pi=3.0\%$ for minimum ionizing pions [64]. Figure 30 shows for a number of gas mixtures the measured separation power at 15 GeV/c, obtained from the lowest 40% pulse heights from 64 samples of 4 cm length [78]. The relativistic rise at 1 atm is highest for noble gases, around 1.6–1.7 with respect to the minimum dE/dx. The low-Z molecular gases show better resolution $\sigma(I)$ but have a lower relativistic rise. There is no 'magic gas' with outstanding PID properties in all respects, although there are significant differences. For the choice of gas in a particular experiment, other characteristics such as diffusion, drift velocity and radiation length will enter, in addition to PID.

For rough estimates of the relative truncated mean resolution $\sigma(I)/I$ as a function of pressure p, sample size d and number N of samples in mixtures of Ar with 10–20% CH₄ the following relation has been derived from the PAI model for pure argon [75]:

$$\sigma(I)/I = 0.41(pd)^{-0.32}N^{-0.46} = 0.41(pL)^{-0.32}N^{-0.14},$$
(3.6)

where L is the total length over which I is measured. To allow comparison of different gases, an extension was proposed:

$$\sigma(I)/I = 0.345(Apd)^{-0.32}N^{-0.46},$$

with
$$A = 6.83 \nu p D/(\beta^2 I)$$
, (3.7)

where ν is the mean number of electrons per molecule. For He, Ne, Ar, Kr and Xe, $A=0.32,\,0.50,\,0.62,\,0.65$ and 0.70, respectively. Values obtained from these theoretical relations based on a likelihood treatment were intended as a guide only to estimate best possible resolutions. Based on experimental results, it has been argued that below a sample size of about 5 cm atm in argon the gain in resolution is negligible [79]. More recent studies, however, indicate that sampling down to 0.5 cm atm still improves the accuracy [80].

Table 4 gives an overview of the PID performance of some TPCs. Included are also the jet chamber of OPAL at LEP and two big detectors, the EPI and ISIS2, which were dedicated to dE/dx measurements with only restricted tracking capability. The best measured values for $\sigma(I)$ are compared with the ideal resolution obtained from the relations cited above, assuming the maximum number $N_{\rm max}$ of samples can be used. The last row shows the calculated resolution for the case that only 70% of $N_{\rm max}$ samples are useful, which is closer to reality in most cases. The measured values are very close to these theoretical numbers. For PEP4 and EPI, the best values are obtained for very clean tracks, where the assumption of $N_{\rm max}$ contributing samples is reasonable.

4. Particularities of some TPCs

4.1. Global aspects

Many TPCs have been constructed. They may be grouped crudely according to the type of experiments they were or are part of.

(a) Electron/positron storage rings: PEP4, TOPAZ, ALEPH and DELPHI.

These experiments demanded large TPCs but had to handle only low rates of events with multiplicities of 2–30 charged tracks. Their designs are similar. PEP4 and TOPAZ were pressurized to maximize PID. ALEPH, the largest of these four TPCs, with the longest drift length (2.2 m) of all TPCs up to now and DELPHI put emphasis on momentum resolution, pattern recognition and reduced material in front of other components and chose operation at 1 atm. The size of the DELPHI TPC had to be reduced because of the addition of a RICH for more powerful PID. Both ALEPH and DELPHI used circular pad rows to improve spatial resolution. ALEPH chose longer pads for better momentum resolution at the highest energies, DELPHI preferred shorter pads for better two-track separation of lower momentum particles.

A reconstructed event from the PEP4 TPC is shown in figure 31.

(b) Heavy ions: fixed-target and collider experiments.

Medium-sized and large TPCs have been used in a number of fixed-target experiments: EOS/HISS at the BEVALEC, the BNL 810 TPC at Brookhaven and NA35, NA36 and NA49 at CERN. A very large TPC is operating for STAR at Brookhaven and an even larger one, ALICE, is ready for data taking at the LHC at CERN. All these TPCs have to handle high particle multiplicities up to several thousand per event in STAR and even more in ALICE. They use(d)

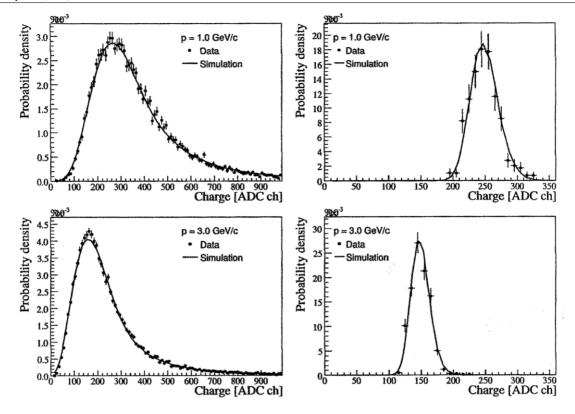


Figure 27. Simulated and measured straggling functions (left) and truncated means (right) for protons [76].

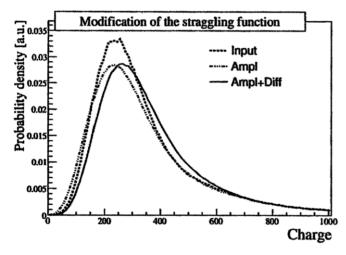


Figure 28. Effect of fluctuations due to amplification and diffusion on the straggling function [76].

1.0 x in Ar cm 1 2 4 8 0.8 cm cm cm $f(\Delta/x)$ 0.6 <∆/x> 0.4 0.2 0.0 1000 2000 3000 4000 Δ/x (eV/cm)

Figure 29. Simulated straggling functions of protons for segments of different lengths x, normalized to x. (Reprinted with permission from [77]. Copyright 2006, Elsevier.)

cathode planes subdivided into a large number of pads, or densely spaced short sense wires (NA36, BNL810).

EOS was an early heavy ion experiment, using full pad coverage. It had a spatial resolution of $300 \,\mu\text{m}$ and a two-track separation of 2.5 cm. The powerful mass separation by dE/dx is demonstrated in figure 32 [81].

NA 49 consisted of four large TPCs, two upstream inside a magnetic field, two outside, providing a long lever arm for momentum measurement. Simple thin field cages were used, in the form of Al-coated Mylar strips stretched over posts, separated from the gas container. The maximum drift length was 1.1 m vertical, pads on the top. Figure 33

gives an indication of the PID in the relativistic-rise region by ionization sampling.

STAR, 4.2 m long and 4 m in diameter, has operated in Au on Au collisions with 200 GeV/nucleon at average luminosities up to 8×10^{27} [83]. The unique pattern recognition of the TPC in multi-track events with more than 3000 tracks per central Au–Au event is visible in the display of a collision, figure 34. STAR was designed to separate pions from protons up to 1.2 GeV/c. Figure 35 shows the energy loss distribution for primary and secondary particles. On a statistical basis, PID has been achieved even in the relativistic-rise region.

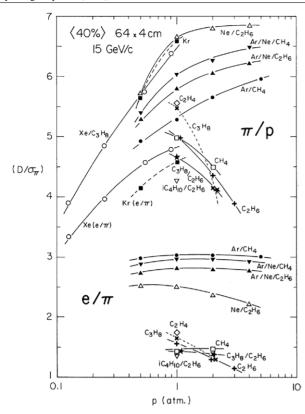


Figure 30. Separation power *D* from ionization sampling, obtained at 15 GeV/*c* with 64 samples of 4 cm. (Reprinted with permission from [78]. Copyright 1982, Elsevier.)

Starting first operation at the LHC, ALICE is the largest TPC constructed so far, with 5 m diameter and 2.5 m drift length. To handle event multiplicities still higher than those of STAR, some 560 000 pads are read out, making use of highly integrated state-of-the-art electronics, see section 3.2.7. Despite the high occupancy, PID is aimed for with a resolution of about 5% [76].

(c) A group of TPCs with rather specific demands. A few of them will be discussed below.

4.2. Special aspects

TRIUMF. The experiment was designed specifically to search for the rare decay $\mu Z \rightarrow e^- Z$ at the level of 10^{-12} [84]. PID was required over a large solid angle surrounding the target with good cosmic ray rejection, momentum resolution of about 1% and tolerance to high rates (10^6 stopping muons/s). The novel TPC idea was picked up and with a simplified design and operation at 1 atm, the experiment could start around the same time as PEP4. The chamber had only 144 anode wires, each sitting in a rectangular cathode with a 6 mm slot and above a pad row with $6 \times 19 \, \mathrm{mm}^2$ pads. The maximum drift length was $34 \, \mathrm{cm}$ in $0.9 \, \mathrm{T}$.

CERES/NA45. CERES [85] is a DC with a radial, variable drift field, 2 m long and 2.6 m in diameter. It is located inside a very inhomogeneous magnetic field formed by two separated short coils. Calibration with laser beams is essential. It has about 15 000 pads of staggered chevron type. A resolution

of 250–350 μm in azimuth and 400–500 μm radially was achieved.

MUNU TPC. The MUNU TPC [86], to study antineutrinoelectron scattering near the Bugey reactor, kept the same structure as the Gotthard TPC (see below) but with a larger fiducial volume of 10^3 L and was filled with CF₄ at 3 atm. The power of continuous pattern recognition is again used to differentiate between electrons from Compton scattering, pair production and ββ-decays. The TPC was contained in an acrylic vessel with fieldshaping rings, and immersed in a tank filled with liquid scintillator, viewed by photomultipliers. Spatial resolution was improved by better electronics but still limited by noise, as well as by granularity. Various R&D activities are pursued in view of the construction of a significantly larger TPC. One interesting result is that Micromegas detectors with a 225 μm gap can achieve a gain above 10^3 in Ar/CF₄ (98/2) at 4 atm.

MuCap. The muon capture experiment at the PSI has used a TPC 15 cm wide, 28 cm long and with a 12 cm drift height, filled with hydrogen at 10 atm and room temperature and operating at a gain of 60 [87]. The TPC allows offline definition of a clear fiducial volume 15 mm away from any material. For the next phase, a modified TPC has been commissioned to operate in D_2 at 30 K and 4 atm and will read pad signals with a unit gain.

HARP. The HARP TPC [88], with 1.6 m drift length and 82 cm diameter and operated in Ar/CH₄ (91/9), has been used at CERN as a large-angle spectrometer, surrounding the target, for precision measurements of hadron production by protons and pions colliding with nuclei, making use of track reconstruction and PID. Corrections have been elaborated, which reduced serious track distortions due to inhomogeneities of the electric and magnetic fields and to beam variations to insignificant level and resulted in good physics performance.

Liquid argon TPC.

- ICARUS. A very large DC filled with liquid argon has been constructed and tested for the ICARUS project [89]. The T600 module contains 600 ton of argon, is 18 m long and has a maximum drift length of 1.5 m. The xy-information is not obtained from pads but from three wire planes 3 mm apart, with wires (150 μ m \varnothing at 3 mm pitch) running at 0° , $+60^{\circ}$ and -60° , by induction when electrons are passing through the first two planes before they are collected on the last plane. The reconstruction of 3D points is based on the fact that the drift time coordinate is shared amongst the three wire planes. The charge deposited by the particle is about 1.5 fC mm⁻¹. As the chamber works without amplification in the liquid, amplifiers with very low noise are required. The purity level of the LAr has to be extremely high, allowing only around 1 ppb of oxygen to keep electron attachment low. Excellent pattern recognition and energy resolution was achieved for cosmic ray events. An example [90] is shown in figure 36.
- ArgoNeuT. The Argon Neutrino Teststand (ArgoNeuT) is installed at Fermilab in the NuMI neutrino beam upstream

Table 4. Best measured resolution values as compared with simulations [75], which assume N_{max} or 0.7 N_{max} as number of samples.

		Experiment					
Param.	PEP4	ALEPH	DELPHI	STAR	OPAL	EPI	ISIS2
$\overline{N_{ ext{max}}}$	183	340	192	45	120	128	320
D (cm)	0.4	0.4	0.4	13×11.5 32×19.5	1.3	6	1.6
p (atm)	8.5	1	1	1	4	1	1
L (cm)	73	136	78	77	160	768	512
Best $\sigma(I)$ meas.	2.7	4.4	5.7	7.4	3.1	2.55	3.1
Ideal resol., N_{max}	2.5	3.8	4.9	6	2.7	2.5	2.5
$N=0.7 N_{\rm max}$		4.4	5.8	7.1	3.1		2.9

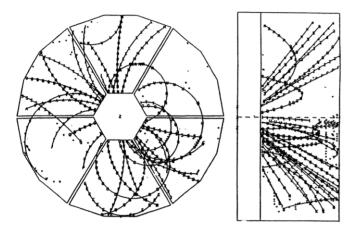


Figure 31. A reconstructed event in the PEP4 TPC; $r\varphi$ and rz views.

of MINOS [91]. Its purpose is to study interactions of low energy neutrinos (0.1–10 GeV) in a LAr TPC and gain valuable technical experience with construction and operation in view of building a similar detector on a much larger scale. Dimensions of the sensitive volume are $47 \times 40 \times 90 \,\mathrm{cm}^3$. The read-out structure is similar to that of ICARUS but with 4 mm wire pitch and wire plane separation. The TPC started operation in 2009.

- MicrobooNe. MicrobooNe [92], an experiment at Fermilab, is to build and operate a LAr TPC with about 100 ton. The experiment will measure low energy neutrino cross-sections and investigate the low energy excess events observed previously, profiting from its good electron—gamma separation. The detector is considered as a necessary next step towards the construction of a kilo ton range detector.
- ArDM. The goal is the operation of a bi-phase LAr detector of about 1 ton for dark matter search, 'with independent ionization and scintillation read-out, to demonstrate the feasibility of a noble gas ton-scale experiment with the required performance to efficiently detect and sufficiently discriminate backgrounds for a successful WIMP detection' [93(a)]. The read-out is based on the Large Electron Multiplier (LEM)-TPC idea. Electrons drifting in the liquid argon are extracted at the liquid-vapour boundary into the vapour volume by the field of a double-grid, with one grid inside the liquid and the other in the vapour. From there they are attracted to and

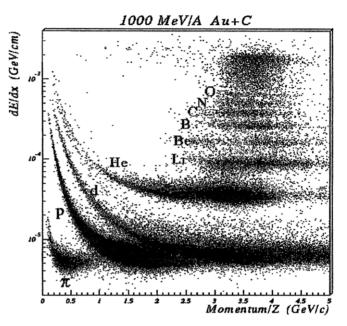


Figure 32. Mass separation by dE/dx in EOS [81].

amplified in two LEMs and captured by an anode above. Signals are read from crossed strips on the anode and upper side of the second LEM. A LEM has a Gas Electron Multiplier (GEM)-like structure (see section 5.2.2) but with larger dimensions: $500~\mu m$ holes on a $800~\mu m$ pitch in a 1.6 mm thick FR4 board. This design is supposed to facilitate large area construction.

A small prototype (10 cm² surface and 10 cm drift in the liquid) has been constructed to test the technique [93(b)]. With cosmic muons, an effective gain of 10 has been achieved. This gain is considered sufficient to compensate the attenuation of charge collection for up to 10 m drift in LAr. On the other hand, gains >100 are required to reach a low enough energy threshold for direct dark matter searches.

Another challenge for ArDM is to obtain an electron/recoil rejection factor $> 10^9$ (required because of the contamination of natural Ar with the isotope 39 Ar, a beta-emitter with $Q = 565 \, \text{keV}$) by measuring very precisely the ratio of scintillation to ionization yields. Various light detection systems are being studied.

 GLACIER. The Giant Liquid Argon Charge Imaging ExpeRiment (GLACIER) is a large underground

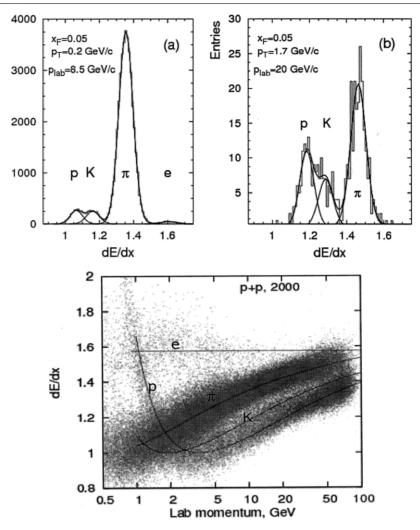


Figure 33. PID by dE/dx in the relativistic-rise region from NA49. (Reprinted with permission from [82]. Copyright 2008, IEEE.)

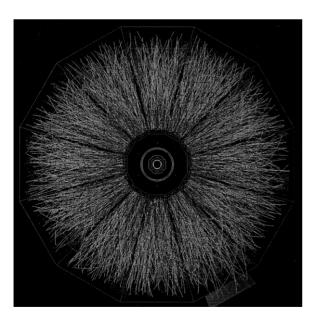


Figure 34. A reconstructed central Au–Au collision in STAR [83(b)].

observatory for proton decay search, neutrino astrophysics and CP-violation studies in the lepton sector [94]. The concept should be scalable up to a total mass of 100 kton. The proposed cylindrical cryostat is based on industrial liquefied natural gas technology. Electrons have to be drifted over 20 m, requiring special care for achieving and maintaining high purity in the LAr and for the drift voltage of the order of a megavolt. The read-out is based on the novel LEM technique (see above). Compared with the 'standard' wire read-out, this technique is considered easier to scale to large size. In GLACIER there is also the possibility to detect Cherenkov light and 'conceiving to have a high $T_{\rm c}$ superconducting solenoid in LAr'.

The project is supposed to proceed in steps, the first one being ArDM. The next step considered is a 10 ton device, to test and optimize the read-out methods and assess the calorimetric performance. Beyond these developments, 'a 1 kton scale device is considered the appropriate choice for a full engineering prototype of a 100 kton detector.'

Xe TPCs. The 136 Xe isotope is a good source for studying neutrinoless double beta decay. The energy released is $Q=2.458\,\mathrm{MeV}$, higher than most of the radioactive decays from

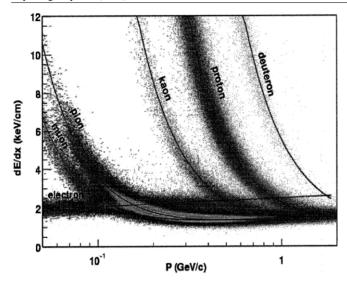


Figure 35. PID in very high-multiplicity events in STAR. (Reprinted with permission from [83(a)]. Copyright 2003, Elsevier.)

Run 975, Event 15 Collection Left

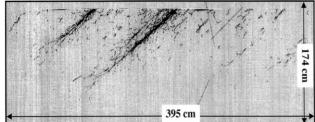


Figure 36. Pattern recognition in the liquid argon TPC of ICARUS. (Reprinted with permission from [90(b)]. Copyright 2003, Elsevier.)

the U and Th chains. Nevertheless, excellent background suppression for energies near Q is necessary to identify or exclude the neutrinoless events. The clear definition of the fiducial volume combined with continuous tracking and good energy resolution of a TPC are crucial for this. Both gaseous and liquid fillings have been proposed. Gas offers in principle better energy resolution, electron tracking with identification of Bragg peak(s) and angular correlation and possibly easier identification of the final state (136 Ba⁺). But the detector is less compact and scaling up is more difficult (pressure vessel).

The Gotthard TPC [86] was filled with $180 \, \text{L}$ of Xe gas at 5 atm (5 kg), enriched to 68% in ^{136}Xe with 5% CH₄ as quencher and used in the Gotthard tunnel. The read-out was at the top with a standard grid and anode/potential wire plane. The cathode plane had perpendicular x and y strips with 4 mm pitch. The anode signal was used to measure the total charge. A double beta decay candidate would show a single continuous track within the fiducial volume with enhanced energy deposition at both ends. In addition, an energy window around $2.46 \, \text{MeV}$ is defined with an energy resolution of 2.7% rms for the neutrinoless decay. No excess above background was found, which set a lower limit of the half-life of 4.4×10^{32} years.

NEXT [95] is another proposal for a TPC using 100 kg of enriched gaseous Xe at 10 bar, in a cylinder about 2 m long and 1 m in diameter. A first stage with 10 kg of natural Xe, Next-10, is foreseen to be built during the next 2–3 years.

Recently, an effort [96] has started to reach unprecedented energy resolution by not only measuring the ionization and scintillation light but by reducing the fluctuations from the amplification of the ionization signal, which destroy the benefit of the small Fano factor in Ar. Some amplification is required, as the fluctuation of the total charge from an absorption of 2,46 MeV is 120 electrons rms and electronic noise would dominate without gas amplification. The idea is to accelerate the electrons to an energy sufficient to excite optical states but not to ionize. The fluctuations from the electroluminescence should be small, as there is no avalanche process involved: photons do not excite additional emissions. A test setup is being assembled at LBNL to demonstrate electroluminescent internal gain in a high pressure gas. Photomultipliers record primary scintillation and provide a start for the drift time, SiPMs perform tracking.

A liquid Xe TPC, *EXO-200*, is presently being commissioned at WIPP [97]. It uses 200 kg of enriched liquid Xe and detects both scintillation and ionization. It is meant as a prototype for a ton-scale detector. Exo-200 is a cylinder, 35 cm long and 40 cm in diameter. It has a central cathode and both ends are equipped with induction and charge collection crossed wire grids followed by large avalanche photodiodes (1.6 cm active diameter). The simultaneous read-out of scintillation and ionizations improves the energy resolution.

A LXe TPC has also been proposed for a small PET system (inner diameter 14 cm, outer 38 cm, length 8 cm), making use of the high light yield and short decay time (98% in 2.2 ns) in Xe and the good spatial and energy resolution of a TPC [98]. A small test chamber gave a combined energy resolution of 3.9% rms from light measured with APDs and from ionization of 511 keV electrons. First tests of a sector prototype ($2\pi/12$) started in 2008. Two sidewalls carry APDs and the read-out is from anode wires and cathode strips.

4.2.1. Radial TPCs.

STAR Forward TPCs. Two cylindrical forward TPCs have been added to the STAR experiment to increase the coverage at small angles to the beam [99]. A radial drift field (perpendicular to the solenoid field) was chosen to improve 2-track separation close to the beampipe to 1–2 mm. Each 1.2 m long TPC is split into five planar slices with an inner tube on HV (R = 7 cm) and an outer wall at ground potential $(R = 30 \,\mathrm{cm})$ and containing the read-out chambers. The chambers follow the curvature and cover $2\pi/6$. They contain two rings of pad rows with pads 1.9 mm in φ and 20 mm along the beam. The anode wires are at 1.5 mm from the pads and cross these at 17.4° to avoid a steplike structure of the response around φ . An Ar/CO₂ (50/50) mixture was chosen for low diffusion and small Lorentz angle. Because of the slow drift, a pulse shaping time of 350 ns was retained and 5 MHz sampling.

BONUS. A small TPC (12 cm diameter, 20 cm length) has recently been operated at Jefferson Lab in studies of neutron structure from electron–neutron interactions [100]. It allows identification and measurement of spectator protons. The radial field was preferred for better acceptance of low-momentum protons. It is the first experiment using a curved

Triple-GEMs for gas amplification. A He/DME (4/10) mixture was chosen to minimize energy loss by the protons and to provide stable operation with sufficient gain.

4.2.2. Spherical TPC. The idea of a spherical TPC is being investigated around various applications demanding detection of low energy neutrinos. The NOSTOS project [101], for the observation of neutrino oscillations, foresees a TPC with an inner sphere of 1 m diameter and an outer sphere of 20 m diameter. The inner sphere should contain some 20 kg of tritium and be covered with Micromegas detectors (see section 5.2.1). The inter-sphere drift volume is to be filled with Xe at 10 atm. The spherical geometry has the advantage of minimizing the detector surface, being a source for background, for a given volume. A prototype spherical DC only sampling radial information (i.e. not providing 3D points) is being tested. The outer vessel has 1.3 m diameter. In the centre is mounted a 1 cm diameter sphere acting as single readout channel. With Ar/CO₂ (90/10) and Ar/Isobutane (98/2) a gas amplification above 10⁴ has been achieved.

This structure has recently been used with an inner sphere of 1.6 cm diameter as a proportional counter. It has shown good energy resolution [101(b)] and has been used to measure thermal neutrons with a ³He filling [101(c)]. A rise time cut permitted good background rejection.

5. Gas amplification without anode wires

To date, gaseous TPCs of almost all experiments had MWPCs as read-out chambers, one exception is the T2K experiment, which started taking data with TPCs using Micromegas read-out. Out of a number of read-out techniques avoiding anode wires, which have been proposed for TPCs, we shall discuss three options: parallel plate chamber (two versions), Micromegas and GEM structures. The first one is a relatively old idea, the latter two have raised interest mainly in connection with TPC projects for a future $0.5-3\,\text{TeV}$ LC. The goal is improved spatial resolution by suppressing the $E\times B$ contribution and finer granularity for two-track separation. The present design specifications for a TPC at the LC are:

1 0 1	
Number of points/track	up to 200-250
Point resolution, rms	100–140 μ m ($r\varphi$);
	\sim 1 mm (z)
Double-track separation	$2-3 \mathrm{mm} (r\varphi); \sim 1 \mathrm{cm} (\mathrm{z})$
Momentum resolution (TPC)	$\sigma/p_{\rm t}^2 \sim 2 \times 10^{-4} { m GeV^{-1}}$
dE/dx resolution, rms	4–5%
Minimum material	$4\% X_0$ centrally, $15\% X_0$
	forward

Thus, the demands for the LC go beyond the performance of existing designs with respect to spatial resolution, two-track separation and material thickness.

5.1. Parallel plate chamber

Parallel plate chamber technology has since long found many applications, in the discharge mode as fast counters and spark chambers and in a 'sub-discharge' mode in streamer chambers. During the final development phase for the LEP

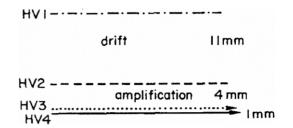


Figure 37. Principle of the parallel plate avalanche chamber. (Reprinted with permission from [102]. Copyright 1983, Elsevier.)

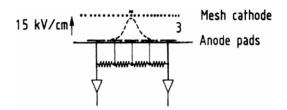


Figure 38. Principle of the pad TPC. The drift volume lies beyond the mesh. (Reprinted with permission from [103]. Copyright 1983, Elsevier.)

TPCs, two proposals were made simultaneously for a use in the proportional mode.

The first proposal [102], called parallel plate avalanche chamber (PPAC), grew out of the development of multigap avalanche chambers and is sketched in figure 37. One coordinate is read out in the HV3 plane from $100 \, \mu \text{m}$ wires with a $500 \, \mu \text{m}$ pitch, the other one from orthogonal wires or pads in the HV4 plane. Tests were done with wires read-out, with and without additional amplification between HV3 and HV4. HV2 is a wire mesh.

The other proposal, called pad TPC [103], grew out of the idea to avoid amplifying wires altogether and to have a natural geometry for implementing circular pad rows, see figure 38. The pads are the anode, the HV being applied to the mesh cathode, which terminates the drift volume. Circular pad rows had at that time not yet been implemented in the standard TPCs with amplifying anode wires. It was thought that complicated corrections would be required due to the variable angle between wire and pad row directions. This finally turned out not to be a problem and circular pad rows were introduced by ALEPH and DELPHI. In the proposal for a pad TPC, resistive charge division had also been suggested, with discrete components or with a resistive layer, to increase the read-out pitch (lowering the channel count and cost) but keeping the good spatial resolution [104].

Despite the fact that none of the proposals was followed up because MWPCs were considered more advanced and thus a safer choice, it is useful to look at the advantages claimed, because these re-appear in the new options presently under study.

Gains up to 10^5 were achieved with Ar/CH₄ mixtures and amplification gaps of 3 and 4 mm. A gain G of 3×10^4 requires an amplification field $E_A = 13.75 \,\mathrm{kV \, cm^{-1}}$ with a 4 mm gap in Ar/CH₄ (80/10). As $G = \mathrm{e}^{\alpha d}$, where α is the first Townsend coefficient and d is the gap width, one deduces $\alpha = 25.8 \,\mathrm{cm^{-1}}$.

• *Positive ion feedback*. As the ion feedback ratio into the drift volume depends on the ratio of the drift field divided

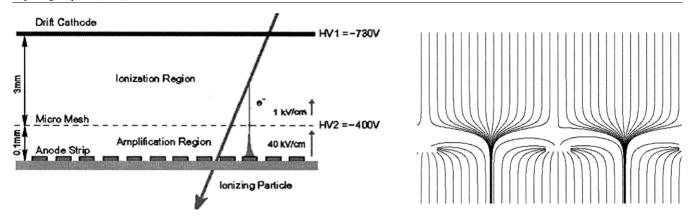


Figure 39. A schematic cut through the Micromegas. (Reprinted with permission from [112]. Copyright 2004, Elsevier.)

by the amplification field it is around 1% in the parallel plate amplification scheme compared with 10–20% in the standard MWPC.

• Two-track separation. The avalanche has a width of only 1 mm FWHM, as measured in the PPAC. A two-track separation in $r\varphi$ down to 2 mm should in principle be achievable. This is much better than with the standard TPC. A read-out with closely spaced, thin pads, however, would be required to make use of this potential, possibly with resistive charge division.

The two-track separation in the drift direction can in principle also be of the order of 2 mm with fast pulse shaping. The fast electron movement induces about one tenth of the total charge in ≤ 10 ns, as most of the charge is produced at a distance of about $(1/\alpha)$ cm. The ions move so slowly that they contribute very little: with integration over 100 ns the total induced charge would be increased only by 10% to 12.5% from the contribution of CH₄-ion drift. Using the electron signal means of course an effective gain of $G_{\rm eff} = {\rm e}^{\alpha d}/\alpha d$ only, i.e. in our example about one tenth of the gas amplification. Pads in a standard TPC with field wires see an effective gain of <20% of the avalanche charge at 1 atm and less at higher pressure.

• Space point resolution. Due to the large suppression of the $E \times B$ effect in the PPAC (with narrow pitch of the anode wires) and almost complete suppression in the pad TPC, a significant improvement on the spatial resolution in $r\varphi$ is expected and the suppression of the asymmetry between positive and negative curvature.

5.2. Micromegas

A schematic view of the Micromegas (MICRO-MEsh GASeous structure) [105] and the electric field lines is shown in figure 39. The electrons from the track ionization pass through a fine mesh into the amplification gap and are multiplied in the high field, inducing signals on the anode plane segmented into read-out pads. The 'micromesh' is woven from $16 \,\mu\text{m}$ wires, leaving holes of about $(50 \,\mu\text{m}^2)$. In contrast to the pad TPC, the Micromegas has a gap of only $(50-100) \,\mu\text{m}$. Small pillars support the mesh every 2.5 mm to guarantee adequate flatness.

Chambers have been in use in some experiments, e.g. in Compass, where 12 chambers $(40 \times 40) \text{ cm}^2$ obtained

resolutions of 70– $90\,\mu m$ and 9 ns and operated in particle fluxes up to 25 MHz mm⁻².

To obtain sufficient amplification in the thin gap, a very high electric field of $30\text{--}80\,\mathrm{kV\,cm^{-1}}$ is required. As a consequence of the high amplification-field/drift-field ratio, the electrons enter through a very narrow funnel. Electron diffusion increases in the high field and causes the avalanche to spread beyond the original funnel to a width of about $35\,\mu\mathrm{m}$ FWHM. This in turn is reducing the positive ion feedback to about 0.2%, as ions produced outside the original funnel will be trapped by the grid. No change of ion feedback was observed for $B = (0-2)\,\mathrm{T}$.

The signal formation in a Micromegas is very fast, also the component due to the positive ions, as these reach the mesh in less than 50–100 ns. As a result, even with fast signal shaping most of the total avalanche charge $N_0 e^{\alpha d}$ is seen and not only the electron component $N_0 e^{\alpha d}/\alpha d$.

For pad widths $\geq (1-2)$ mm, the avalanche is too narrow and the 'pad response width' has to be broadened to enable charge interpolation. This broadening has been achieved by laminating to the anode a Mylar film coated with a high surface resistivity material (e.g. CERMET-Al-Si alloy with 1 M Ω /square). With pads of 2 × 6 mm and a gap of 50 μ m, a resolution of $50 \,\mu m$ was measured with gas gains of 2300 and 4700, not depending on drift length for up to 15 cm drift in 5 T. A gas mixture of Ar/CF₄/isobutane (95/3/2), the so-called T2K gas, was chosen for its high electron drift velocity $(7.3 \text{ cm } \mu\text{s}^{-1})$ at $E = 200 \text{ V cm}^{-1}$ and for the high $\omega \tau$ at 5 T, reducing the single-electron transverse diffusion to $190 \,\mu\text{m}/1 \,\text{m}$ drift [106]. From the measured resolution it was estimated that some 27 electrons (= N_{eff}) were statistically relevant for the centroid determination. This is close to the most probable ionization for 6 mm long pads.

Also the Micromegas has some $E \times B$ effect in the holes of the grid but this is insignificant because of the small dimensions.

Various systems are being studied, in which the printed circuit board anode with pads is replaced by Si-pixel devices ('Digital TPC'), with and without enlarged metallic pixels, with a separate Micromegas mesh or an integrated grid and with resistive protection layers of a-Si: H or Si₃N₄ [107]. The digital pixel read-out does not require signal broadening.

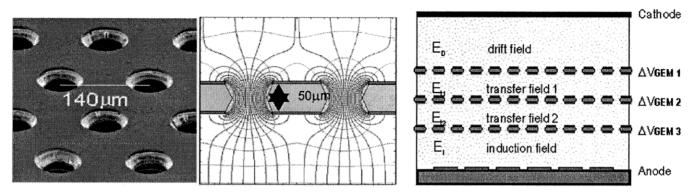


Figure 40. Principle of the GEM structure. (Reprinted with permission from [112]. Copyright 2004, Elsevier.)

The first large experiment using Micromegas as read-out chamber for a TPC is T2K [108]. It employs what is called 'bulk-Micromegas', in which the whole structure combining the multilayer read-out plane, the mesh and the pillars produced by photolithographic etching procedures as a unit, in form of (40×40) cm² plates [109]. The gap is $128 \,\mu\text{m}$ and the 0.3 mm pillars have a pitch of 2.5 mm, providing about 93% active surface. The pads are about (7×10) mm², read out via blind holes and screened lines on the back. A gain variation of 2.2% was achieved after corrections for capacitance. Operating in 0.2 T, the TPCs are to provide a 10% momentum resolution at 1 GeV/c and separation of electrons from muons by dE/dx.

MIMAC [110] is a multi-chamber detector for dark matter search. It is planned to do tracking and measure ionization with a matrix of TPCs filled with 3 He and CF₄ (gases with sensitivity to spin-dependent interactions), with Micromegas as detector with high granularity read-out. As a first step, a small prototype has provided the first measurement of He ionization in He down to recoil energies of 1 keV, important for the use of He in dark matter search. In a second step, 3D reconstruction of tracks of 5 MeV α-particles has been achieved in He/isobutane (95/5) at 800 mbar. For this measurement the anode had x and y strips at a 200 μm pitch. The tests are being extended to recoil energies below 6 keV, the range of interest in dark matter search.

5.3. Gas electron multiplier

The GEM [111] is produced by chemically etching holes at high density into a thin metal-coated polymer foil. Gas amplification is achieved inside the holes by applying a voltage on the metal layers. Figure 40 shows the layout of the holes, electric field lines and a 'triple-GEM' structure with an anode plane for read-out [112]. Typical parameters are foil thickness = $50 \, \mu \text{m}$, hole pitch = $140 \, \mu \text{m}$, inner hole diameter = $70 \, \mu \text{m}$, voltage = around $400 \, \text{V}$.

Three GEMs are usually put in series, to obtain sufficient gain with acceptable discharge probability. From the last GEM foil, the electrons drift through the induction gap to the anode segmented into pads or strips. For homogeneity reasons, the minimum gap width is about 1 mm. The induced pulse has an intrinsic spread of about 350 μ m rms. The signal contains basically only a fast electron component, as there are no ions produced in the last induction gap except very near the holes of the last GEM foil.

A system of 20 triple-GEMs with an active area of (31 \times 31) cm² was operated in COMPASS [113], in a high intensity muon beam. A spatial resolution of 70 μ m was achieved at rates up to 2.5 MHz cm⁻². With 50 ns pulse shaping and an effective gain of 8000, the efficiency was 99%. The much shorter peaking time of 10 ns applied in LHCb [114] led to a lower efficiency of \geq 96% for two triple-GEMs in OR. The gain was 6000 in Ar/CO₂/CF₄ (45/15/40).

Several experiments have observed charge build-up on the insulator inside the holes, which changes with particle flux and causes gain changes, reproduced by simulation [115].

For the TPC development, various read-out schemes are being studied, similar to those with Micromegas: standard printed circuit boards with pads, as well as Si pixels. High resolution ($40 \mu m$ for short drift) and granularity was demonstrated with a combination of a triple-GEM and Medipix/Timepix read-out with (55×55) μm^2 pixels [116].

Ion feedback is intrinsically low in a triple-GEM and further reduced in a magnetic field, to 0.25% at $B=5\,\mathrm{T}$ with alternating low and high drift/transfer fields [117(a)]. Similar values are achieved with equal transfer fields but very low drift field ($<100\,\mathrm{V}\,\mathrm{cm}^{-1}\,\mathrm{atm}^{-1}$) in the TPC [117(b)].

The LC-TPC collaboration has constructed a TPC prototype, 60 cm long and 70 cm in diameter, to study many aspects of a TPC for the LC, including various Micromegas and GEM read-out structures [118]. Some of the developments mentioned are certainly very encouraging but most are still far from use on a large scale.

6. Conclusions

Designed some 30 years ago to provide complete event information at low event rates around 1 Hz and with particle multiplicities around 20 particles per event, TPCs have developed into unique devices to study heavy ion collisions with extreme multiplicities of up to several thousand particles per event and registering some hundred events per second. There is no competitor in this field delivering similar event details on a large scale. Besides this, TPCs have found applications in other fields, in particular for the detection of rare reactions/decays. As may be judged from the ongoing projects, it is unlikely that interest will be fading in the near future, because of the unique capability of the TPC to provide

practically continuous 3D spatial and ionization information of high granularity.

References

- [1] Charpak G *et al* 1968 The use of multiwire proportional counters to select and localize charged particles *Nucl. Instrum. Methods* **62** 262
- [2] Bressani T, Charpak G, Rahm D and Zupancic C 1969 Track localization by means of a drift chamber *Proc. of the Int.* Seminar on Filmless Spark and Streamer Chambers (Dubna, Russian Federation) p 275
- [3] Walenta A H, Heintze J and Schürlein B 1971 The multiwire drift chamber—a novel type of proportional wire chamber Nucl. Instrum. Methods 92 373
- [4] (a) Nygren D, PEP-198-1975
 (b) 1976 Proposal for a PEP Facility based on the TPC, PEP4-December 30
- [5] Lohse T and Witzeling W 1992 The TPC The Advanced Series on Directions in High Energy Physics vol 9, ed F Sauli (Singapore: World Scientific) p 82
- [6] (a) CERN Courier January/February 2004
 (b) Lepeltier V 2007 J. Phys. Conf. Ser. 65 012001
 (c) Attié D 2009 Nucl. Instrum. Methods A 598 89–93
- [7] Blum W, Riegler W and Rolandi L 2008 Particle Detection with Drift Chambers 2nd edn (Berlin: Springer)
- [8] Spieler H 2005 Semiconductor Detector Systems (Oxford: Oxford University Press)
- [9] Smirnov I, Heed: Interactions of particles with gases, http://consult.cern.ch/writeup/heed/
- [10] Veenhof R, Garfield–Simulation of gaseous detectors, http://consult.cern.ch/writeup/garfield/
- [11] Biagi S, Magboltz: Transport of electrons in gas mixtures, http://consult.cern.ch/writeup/magboltz/
- [12] (a) Bichsel H 2006 Nucl. Instrum. Methods A 562 154
 (b) Bichsel H 2008 Phys. Lett. B 667 267
- [13] Lapique F and Piuz F 1980 *Nucl. Instrum. Methods* 175 297–318
- [14] Fischle H, Heintze J and Schmidt B 1991 Nucl. Instrum. Methods 301 297
- [15] Amsler C et al (Particle Data Group) 2008 Review of Particle Physics Phys. Lett. B **667** 1
- [16] 1979 ICRU Report 31, Average energy to produce an ion pair (Washington DC: ICRU)
- [17] Lehraus I et al 1978 Nucl. Instrum. Methods 153 343
- [18] Allison W W M et al 1984 Nucl. Instrum. Methods 224 396
- [19] Walenta A H et al 1979 Nucl. Instrum. Methods 161 45
- [20] Blum W et al 1986 Nucl. Instrum. Methods A 252 407-12
- [21] (a) Kobayashi M 2006 Nucl. Instrum. Methods A 562 136–40
 (b) Arogancia D C et al 2009 Nucl. Instrum. Methods 602 403–14
- [22] (a) Pitchford L C and Phelps A V 1982 *Phys. Rev.* A 25 540
 (b) Robson R E and Ness K F 1986 *Phys. Rev.* A 33 2068
 (c) Ness K F 1995 *Aust. J. Phys.* 48 557
- [23] Biagi S F 1989 Nucl. Instrum. Methods A 283 716
- [24] Schmidt B 1986 Doctoral Thesis University of Heidelberg
- [25] (a) Schultz G 1979 Doctoral *Thesis* University of Louis-Pasteur, Strasbourg
 - (b) Schultz G, Chorpak G and Souli F 1977 *Rev. Phys. Appl.* 12 67
- [26] Sauli F 1977 CERN 77-09
- [27] McDaniel E W and Mason E A 1973 *The Mobility and Diffusion of Ions in Gases* (New York: Wiley)
- [28] Hornbeck J H 1951 Phys. Rev. 84 615
- [29] Wagner E B, Davies F J and Hurst F J 1967 J. Chem. Phys. 47 3138
- [30] Schmidt B 1980 Diploma Thesis University of Heidelberg
- [31] 1994 RD-32, CERN/DRDC 94-10

- [32] Amendolia S R et al 1986 Nucl. Instrum. Methods 244 516
- [33] (a) Ishikawa A, http://www-hep.phys.saga-u.ac.jp/ILC-TPC/gas/
 - (b) Ishikawa A, http://www-hep.phys.ac.jp/ILC-TPC/gas/ps/
- [34] Massey H S W, Burhop E H S and Gilbody H B 1969 Electronic and Ionic Impact Phenomena (Oxford: Clarendon)
- [35] McCorkle D L, Christophorou L G and Hunter S R 1981 Proc. 2nd Int. Swarm Seminar (Oak Ridge, TN) (Oxford: Pergamon) p 21
- [36] Bloch F and Bradbury N E 1935 Phys. Rev. 48 689
- [37] Erskine G A 1972 Nucl. Instrum. Methods 105 565
- [38] Ramo S 1939 Proc. IRE 27 584
- [39] Shockley W 1938 J. Appl. Phys. 9 635
- [40] Collected by von Engel A 1956 Handbuch der Physik vol 21 (Berlin: Springer)
- [41] (a) Biagi S F 1989 Nucl. Instrum. Methods A 283 716
 (b) Armitage J C 1988 Nucl. Instrum. Methods A 271 588
- [42] Rose M E and Korff S A 1941 Phys. Rev. 59 850
- [43] Diethorn W 1956 USAEC Report NY6628
- [44] Alkahazov G D 1970 Nucl. Instrum. Methods 89 155
- [45] Curran S C, Cockroft A L and Angus J 1949 Phil. Mag. 40 929
- [46] Legler W 1961 Z. Naturforsch. a 16 253
- [47] Schlumbohm H 1958 Z. Phys. 151 563
- [48] Brand C et al 1985 Nucl. Instrum. Methods A 237 501
- [49] Anderhub M, Devereux M J and Seiler P G 1979 Nucl. Instrum. Methods 166 581–2
- [50] Bourotte J and Sadoulet B 1980 Nucl. Instrum. Methods 173 463
- [51] Hilke H J 1980 Nucl. Instrum. Methods 174 145
- [52] Towrie M et al 1986 J. Phys. B 19 1989
- [53] Dysdale S L T et al 1986 Nucl. Instrum. Methods A 252 521
- [54] Hubricht G et al 1985 Nucl. Instrum. Methods 228 327
 Hubricht G et al 1986 Nucl. Instrum. Methods 243 495
- [55] Hilke H J 1986 Nucl. Instrum. Methods A 252 169
- [56] Gushchin E M, Lebedev A N and Somov S V 1985 Instrum. Exp. Tech. USSR 28 1
- [57] Abele J 2003 Nucl. Instrum. Methods A 499 692
- [58] Lebedev A 2002 Nucl. Instrum. Methods A 478 163
- [59] (a) Va'vra J 1986 Nucl. Instrum. Methods A 252 547
 (b) Va'vra J 2003 Nucl. Instrum. Methods A 515 1
 - (b) Va Vra J 2003 Nucl. Instrum. Methods A 515 1
 - (c) Kadyk J 1991 Nucl. Instrum. Methods A 300 436
 - (d) Tesch N 2002 IEEE Trans. Nucl. Sci. 49 1609
 - (e) Capeans M 2003 Nucl. Instrum. Methods. A 515 73
- (f) Bouclier R et al 1994 Nucl. Instrum. Methods A 350 464
- [60] (a) Kadyk J (ed) 1986 Proc. the Workshop on Radiation Damage to Wire Chambers (Berkeley, CA) LBL 21170
 - (b) 2001 Proc. the Int. Workshop on Aging Phenomena in Gaseous Detectors, (Hamburg, Germany) 2003 Nucl. Instrum Methods A 515 1–386
 - (c) NASA, a very large database on outgassing of materials can be reached via the NASA homepage
- [61] Adam J et al 1983 Nucl. Instrum Methods 217 291
- [62] Malter L 1936 Phys. Rev. 50 48
- [63] ATLAS Collaboration 2008 The ATLAS Experiment at the CERN LHC J. Instrum. 3 S08002
- [64] Lynch G R1987 Performance of the PEP4 TPC Seminar at MPI-Munich (Munich, Germany, 22 June 1987)
 Lynch G R 1991 private communication
- [65] Veenhof R, http://rjd.web.cern.ch/rjd/Harp/charge.html
- [66] Amendolia S R et al 1985 Nucl. Instrum. Methods A 234 47
- [67] Nemethy P et al 1983 Nucl. Instrum. Methods A 212 273
- [68] (a) Jared R C, Landis D A and Goulding F S 1982 IEEE Trans. Nucl. Sci. NS-29 57
 - (b) Aihara H et al 1983 IEEE Trans. Nucl. Sci. NS-30 162
- [69] (a) Campagnolo R 2005 Int. Europhysics Conf. on High Energy Physics EPS (Lisbon, Portugal)
 - (b) Mota B et al 2004 Nucl. Instrum. Methods A 535 500

- [70] Veenhof R, http://rjd.web.cern.ch/rjd/Harp/signal.html
- [71] (a) Decamp D et al 1990 Nucl. Instrum. Methods A 294 121
 - (b) Atwood W B et al 1991 Nucl. Instrum. Methods A 306 446
- [72] (a) Aarnio P et al 1991 Nucl. Instrum. Methods A 303 233
 - (b) Sacquin Y 1992 Nucl. Instrum. Methods A 323 209
 - (c) Abreu P et al 1996 Nucl. Instrum. Methods A 378 57
- [73] Blum W et al 1986 Nucl. Instrum. Methods A 252 407
- [74] Alikhanov A et al 1956 Proc. CERN Symp. On High Energy Accelerators and Pion Physics vol 2, p 87
- [75] Allison W W M and Cobb J H 1980 Annu. Rev. Nucl. Part. Sci. 30 253
- [76] Christiansen P and Baechler J 2006 ALICE-INT-2005-030
- [77] Bichsel H 2006 Nucl. Instrum. Methods A 562 154
- [78] Lehraus I, Matthewson R and Tejessy W 1982 Nucl. Instrum. Methods 200 199
- [79] (a) Walenta A H et al 1979 Nucl. Instrum. Methods 161 45
 - (b) Lehraus I et al 1982 Nucl. Instrum. Methods 196 361
- [80] Assmann R 1990 Diploma Thesis University of Munich
- [81] Ronan M 2004 Second 'rare event' TPC Workshop (Paris, France, December 2004)
- [82] Varga D for NA49 2008 Experiment *IEEE Nuclear Science Symp. (Dresden, Germany)*
- [83] (a) Anderson M et al 2003 Nucl. Instrum. Methods A **499** 659–78
 - (b) http://www.star.bnl.gov/STAR/magelib/collisions2001/ev_2front.jpg
- [84] (a) Hargrove C K et al 1984 Nucl. Instrum. Methods 219 461–71
 - (b) Mes H et al 1985 Nucl. Instrum. Methods 225 547–9
- [85] Adamova D et al 2008 Nucl. Instrum. Methods A 593 203
- [86] Vuilleumier J L 2009 J. Phys. Conf. Ser. 179 012013
- [87] Kammel P for the MuCap Collab 2008 Few Body Syst. 44 333–6 DOI 10.1007/s00601-008-0321-8
- [88] Ammosov V et al 2008 Nucl. Instrum. Methods A 588 294–317
- [89] Rubbia C 1977 The liquid argon time projection chamber: a new concept for neutrino detectors EP Internal Report 77-8, CERN
- [90] (a) ICARUS Collaboration 2004 Nucl. Instrum. Methods. A 527 329
 - (b) Calligarich E 2003 ICARUS, *Time Projection Chamber Symp. (LBNL, Berkeley, CA, October 2003)*
- [91] http://t962.fnal.gov/
- [92] http://www-microboone.fnal.gov/
- [93] (a) http://neutrino.ethz.ch/ArDM/
 - (b) Badertscher A et al 2009 Nucl. Instrum. Methods A 617 188–92
 - (c) Bondar A et al 2008 Nucl. Instrum. Methods A (arXiv:0807.0530)
- [94] Badertscher A et al 2009 arXiv:1001.0076v1[physics.ins-det]
- [95] NEXT: Díaz J et al 2009 J. Phys. Conf. Ser. 179 012005
- [96] (a) Nygren D 2003 TPC Symp. (LBNL, Berkeley, CA, October 2003)
 - (b) Nygren D 2007 Nucl. Instrum. Methods A 603 337
 - (c) Spieler H et al 2010 VCI Nucl. Instrum. Methods at press
- [97] Gornea R, for EXO Collab. 2009 *J. Phys. Conf. Ser.* **179** 012004
- [98] Amaudruz P et al 2008 IEEE Nuclear Science and Medical Imaging Conf. (Dresden, Germany)
- [99] Ackermann K H et al 2003 Nucl. Instrum. Methods A 499 713–9
- [100] Fenker H et al 2008 Nucl. Instrum. Methods A 592 273–86
- [101] Gorodetzky P et al 2006 Nucl. Phys. B 151 410

- (b) as proportional counter only: Giomataris Y *et al* 2008 *J. Instrum.* **3** P09007
- (c) Andriamonje S *et al* 2009 *J. Phys. Conf. Ser.* **179** 012003
- [102] Peisert A 1983 Nucl. Instrum. Methods 217 229
- [103] Hilke H J 1983 Nucl. Instrum. Methods 217 189
- [104] based on suggestions by Radeka V; (a) Radeka V and Boie R A 1980 Nucl. Instrum. Methods 178 543
 (b) Yu B et al 1992 Nucl. Instrum. Methods A 323 413
- [105] (a) Giomataris Y et al 1996 Nucl. Instrum. Methods A 376 29
 - (b) Giomataris Y 1998 *Nucl. Instrum. Methods* A **419** 239
- [106] Dixit M et al 2007 Nucl. Instrum. Methods A 581 254
- [107] Timmermans J 2009 CLIC09 Workshop (CERN Geneva)
- [108] Giganti C 2009 4th Symp. on Large TPCs for Low Energy Rare Event Detection (Paris, France)
 Giganti C 2009 J. Phys. Conf. Ser. 179 012002
- [109] Giomataris Y et al 2006 Nucl. Instrum. Methods A 560 405
 - de Oliveira R 2009 private communication
- [110] http://lpsc.in2p3.fr/mimac
- [111] Sauli F 1997 Nucl. Instrum. Methods A 386 531
- [112] Hoch M 2004 Nucl. Instrum. Methods A 535 1
- [113] Ketzer B et al 2004 Nucl. Instrum. Methods A 535 314
- [114] LHCb Collaboration 2008 The LHCb Detector at the LHC *J. Instrum.* **3** S08004
- [115] Tikhonov V and Veenhof R 2002 Nucl. Instrum. Methods A 478 452
- [116] Bamberger A et al 2007 Nucl. Instrum. Methods A 581 274
- [117] (a) Killenberg M et al 2004 Nucl. Instrum. Methods A 530 251–7
 - (b) Breskin A et al 2002 Nucl. Instrum. Methods A 478 225–9
- [118] Dehmelt K 2009 TIPP09 (KEK, Tsuba, Japan)
- [119] 1995 IAEA-TECDOC-799 Atomic and Molecular Data for Radiotherapy and Radiation Research (Vienna, Austria: IAEA Nuclear Data Section)
- [120] Gatti E et al 1979 Nucl. Instrum. Methods 163 83-92
- [121] Bellazzini R et al 1985 IEEE Trans. Nucl. Sci. NS-32 389-92
- [122] Fanscher D and Schaffer A C 1979 *IEEE Trans. Nucl. Sci.* NS-26 150

For convenience, references for some TPCs have been grouped together below under the name of the experiment

- [PEP4]: (a) Nygren D R 1974 Proposal to investigate a novel concept in particle detection *LBL Internal Report*, Berkeley
- (b) Proposal for a PEP Facility based on the Time Projection Chamber, PEP4, December 1976
- (c) Fancher D et al 1979 Nucl. Instrum. Methods 161 383-90
- (d) Aihara H et al 1983 IEEE Trans. Nucl. Sci. NS-30 63, 76 and 162
- (e) Cowan G D 1988 PhD Thesis LBL-24715
- (f) Lynch G 1987 Seminar at MPI-Munich (Munich, Germany, 22 lune)
- (g) Jared R C, Landis D A and Goulding F S 1982 *IEEE Trans. Nucl. Sci.* NS-29 57
- [TRIUMF]: (a) Hargrove C K et al 1984 Nucl. Instrum. Methods 219 461
- (b) gating: Mes H et al 1984 Nucl. Instrum. Methods 225 547
- (c) gating: Bryman D A 1985 Nucl. Instrum. Methods A 234 42
- [TOPAZ]: (a) Shirahashi A et al 1988 IEEE Trans. Nucl. Sci. NS-35 414
- (b) Kamae T et al 1989 Nucl. Instrum. Methods A 252 423

- [ALEPH]: (a) Atwood W B et al 1991 Nucl. Instrum. Methods A 306 446
- (b) Decamp D et al 1990 Nucl. Instrum. Methods A 294 121
- (c) gating: Amendolia S R et al 1986 Nucl. Instrum. Methods A 252 403
- [DELPHI]: (a) Aarnio P et al 1991 Nucl. Instrum. Methods A 303 233
- (b) Sacquin Y 1992 Nucl. Instrum. Methods A 323 209
- (c) Abreu P et al 1996 Nucl. Instrum. Methods A 378 57
- [NA35]: (a) The NA35 Collaboration 1989 CERN-SPSC-89-17 and SPSC/M-440
- (b) Jacobs P (NA35 and NA49 Collaborations) 1993 LBL-33810
- (c) Gazdzicki M (NA35 Collaboration) 1995 Nucl. Phys. A 590 197
- [EOS/HISS]: Rai G 1990 IEEE Trans. Nucl. Sci. 37 56

- [NA 49]: (a) Afanasiev S et al 1999 Nucl. Instrum. Methods. A ${\bf 430}$ 210
- (b) Alt C et al 2006 Eur. Phys. J. C 45 343
- [CERES/NA45]: Adamova D 2008 Nucl. Instrum. Methods A 593 203
- [STAR]: Anderson M et al 2003 Nucl. Instrum. Methods A 499 659
- [HARP]: Ammosov V et al 2008 Nucl. Instrum. Methods A 588 294
- [ALICE]: (a) ALICE Collaboration 2008 The ALICE Experiment *J. Instrum.* **3** S08002
- (b) Campagnolo R, HEP 2005 Int. Europhysics Conf., EPS (Lisbon, Portugal)
- (c) Mota B et al 2004 Nucl. Instrum. Methods A 535 500
- (d) Alme J et al 2010 arXiv:1001.1950v1