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# MEASUREMENT OF THE ELECTROSTATIC CHARGE IN AIRBORNE PARTICLES: II – PARTICLE CHARGE DISTRIBUTION OF DIFFERENT AEROSOLS

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**Abstract** - This work gives sequence to the study on the measurement of the electrostatic charges in aerosols. The particle charge classifier developed for this purpose and presented in the previous paper (Marra and Coury, 2000) has been used here to measure the particle charge distribution of a number of different aerosols. The charges acquired by the particles were naturally derived from the aerosol generation procedure itself. Two types of aerosol generators were used: the vibrating orifice generator and turntable Venturi plate generator. In the vibrating orifice generator, mono-dispersed particles were generated by a solution of water/ethanol/methylene blue, while in the rotating plate generator, six different materials were utilized. The results showed no clear dependence between electric charge and particle diameter for the mono-dispersed aerosol. However, for the poly-dispersed aerosols, a linear dependence between particle size and charge could be noticed.

Keywords: Electrostatic charges; Charges measurement; Aerosol particles.

# **INTRODUCTION**

Charged aerosol particles have important implications in the aerosol science due to the electrostatic effects in powder production, handling, transport and sampling (Lui & Pui, 1985) and also on the deposition of respirable particles in the human respiratory tract (Vincent, 1995; Johnston et al., 1985). More particularly, a number of works have demonstrated the importance of electrostatic charges in gas cleaning, as it affects particle collection by filters (Coury, 1983; Duarte Filho, 1985; Coury et al. 1991; Duarte Filho et al, 2000).

In order to investigate electrical effects in aerosols studies, information about the magnitude

and distribution of the particle charge is often needed. For this reason, an improved charge classifier (electrical charge classifier - ECC) was designed and built to measure the absolute particle charge magnitude and distribution. These steps were presented in the first paper by Marra and Coury (2000).

For the current research, a number of dispersed solid particles (e.g. phosphate rock, alumina, dry egg white, wood dust, pyrochlore, diatomite and methylene blue) were used as test powders, and had their charge measured. Two types of aerosol generators have been used: a turntable Venturi generator (TSI Inc., model 3433), for poly-disperse aerosols (phosphate rock, alumina, dry egg white,

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wood dust, pyrochlore and diatomite) and a vibrating orifice aerosol generator (TSI Inc., model 3450), for mono-disperse aerosol (methylene blue).

It was noticed that, during particle generation and dispersion, aerosol particles acquired considerable levels of electrical charge and they were of different electrical signs, and these characteristics varied depending on the powder material and the aerosol generation technique utilized (Marra Jr. *et al.*, 2005). The present study compares the different types of materials with respect to their acquired charge distribution and sign.

## **EXPERIMENTAL SETUP AND PROCEDURES**

A full description of the apparatus and techniques used for the measurement of the distribution, magnitude and polarity of charge carried by the airborne particles is given elsewhere (Coury, 1983; Coury at al., 1991; Marra Jr., 2000; Marra Jr. and Coury, 2000) and only a brief account is given here.

#### **Basic Theory**

A charged aerosol particle will experience an electrical force,  $\mathbf{F}_{e}$ , during transport through a region under an electrical field and its trajectory can be altered, as observed in Figure 1. The electric force,

**F**<sub>e</sub>, is given by:

$$\mathbf{F}_{\mathbf{e}} = \mathbf{Q} \, \mathbf{E} \tag{1}$$

where Q is the particle charge and E is the applied electric field. As a result of this electric force, the particle moves at velocity  $U_x$  in the direction of the line forces, as shown in the Figure 1. The drag force,  $F_d$ , of the fluid on the particle in the direction x can be expressed by:

$$\mathbf{F_d} = \frac{3\mu\pi d_p \mathbf{U}_x}{F_S} \tag{2}$$

where  $\mu$  is the gas viscosity,  $U_x$  is the particle velocity in the x direction,  $d_p$  is the particle diameter and  $F_s$  the Cunningham slip factor.

Assuming that  $\mathbf{F}_{e} = \mathbf{F}_{d}$ , the particle charge can be written as:

$$Q = \frac{3\pi\mu d_p U_0 XL}{Z\Delta VF_S}$$
(3)

where  $U_0$  is the gas velocity, Z is the length of deflection plates, L is the separation distance between the plates and  $\Delta V$  is a potential difference applied between them. Equation 3 is valid to all particles crossing all the electrical field lines, in the direction Z, and to an established velocity profile.



Figure 1: Deflection of a charged particle under an electric field between parallel plates.

# **Electrical Charge Classifier**

The electrical charge classifier, ECC, was built in transparent acrylic and is shown schematically in Figure 2. The apparatus can be divided in to three sections: A – aerosol insertion and gas flow stabilization section, B – particle deflection section, and C – particle collection section. The present design provides a flat profile for the gas velocity in all the extent of region B, with  $U_0=$ 0.30±0.04 m/s (see Marra and Coury, 2000). The classifier is placed in a horizontal orientation with the test aerosol introduced from the central slit in section A.

Once introduced in the classifier, the particle crosses section B and suffers a deflection in its path. The migration of each particle between plates is proportional to its charge and to the applied deflection voltage. The particle is collected in section C, where its deflection and size are measured. Finally, the particle charge is calculated with the use of Equation 3.



Figure 2: General view of the Electrical Charge Classifier (ECC).

#### **Aerosol Generators**

A vibrating orifice generator (TSI Inc. model 3450), capable of producing aerosols from materials in the form of solution, was used to generate a monodispersed aerosol. A solution of water/ethanol/methylene blue was used here. The generation begins with the solution being injected through a small orifice, producing a liquid jet. The orifice is fixed on a device with a piezoelectric crystal, which produces high frequency vibrations when subjected to an electric current. Those vibrations "break" the liquid jet in droplets that are dragged by an air flow, and conducted though a vertical column. In this column, the liquid portion of the droplet evaporates, remaining the non-volatile solute as a solid particle with a uniform and nearly spherical form.

For poly-dispersed aerosol production, a TSI Powder Disperser, model 3433 (DP-3433) was utilized. This is an instrument utilized to disperse small quantities of dry powder with diameters between 1 and 50  $\mu$ m and uses the Venturi aspiration technique. The powder is spread over the surface of a rotating disc with the use of a brush, forming a uniform layer, which is then aspirated by a Venturi through a capillary tube. The lower and upper

sections of the capillary tube are located very close to the surface of the rotating disc and in the narrow portion of the Venturi, respectively. A low-pressure region is created by the air velocity increase in the narrow portion of the Venturi and, consequently, the particles are aspirated through the capillary tube.

# **Test Aerosols**

Methylene blue, with a density of 1.57 g/cm<sup>3</sup> was used in the mono-dispersed aerosol generator, fed as solution in water and ethanol alcohol. After atomization and drying, it resulted in monodispersed particles with the diameter adjusted to the desired value.

Six particulate materials were utilized as polydispersed samples in the rotating plate generator. The materials were chosen with the idea of covering a wide range of origins, compositions and physical/chemical properties. The material properties important in this study (density and median Stokes diameter) are show in Table 1. The densities were measured in a helium pycnometer from Micromeritics Accupyc, Model 1330 and the particle diameters were obtained in a TSI Aerodynamic Particle Sizer (APS), Model 3320.

Material	Density [kg/m <sup>3</sup> ]	Median Stokes diameter, d <sub>pS</sub> [µm]	Geom. S. Dev., $\sigma_g$
Phosphate rock	3,030	1.95	0.60
Alumina	3,698	1.84	0.49
Dry egg white	1,043	3.15	1.36
Wood dust	1,420	3.04	1.04
Pyrochlore	4,110	1.98	0.65
Diatomite	3,860	1.96	0.68

Table 1: Properties of the test aerosols

# **Experimental System**

The complete experimental apparatus used in the analysis of the charged particle behavior can be seen in Figure 3. It can be observed that the ECC was directly connected to the powder dispensers used (one at a time). The dust-laden gas of each test was introduced in the ECC immediately after dispersion. Therefore, the electrostatic charges measured in the tests were those generated in the dispersion technique itself. The particles diameters, after deflection, were measured using the TSI-APS, Model 3320 particle counter.



Figure 3: Schematic view of the experimental unit.

## RESULTS

#### **Charge Polarity**

The charge polarity of the aerosol particles is determined by the peak of their deflection curve within the ECC: if this peak is formed by deflection towards the positive electrode, the particles are predominantly negative, and vice-versa.

Figures 4 and 5 show some typical results of the displacement of charged particles in the deflection zone of the ECC. Figure 4 is relative to 1.93  $\mu$ m particles of pyroclore, whilst Figure 5 refers to 4.71  $\mu$ m particles of alumina. Each curve plots the total of collected particles for a specific diameter d<sub>p</sub>, as a function of the position x, where x=0 means no deflection. These curves were adjusted to the Gaussian distribution. The displacements of charged

particles are proportional to the level of the charging and to the intensity of the applied electric field. A 10 kV potential difference was applied in all tests, with the negative electrode situated at the right side (relative to the aerosol flow) of the ECC.

It can be noticed in Figure 4 that the displacement peak of the 1.93  $\mu$ m pyroclore was positioned to the left side, indicating negatively charged particles. The same behavior was observed for the phosphate rock, the wood dust and the diatomite. An opposite behavior was observed for dry egg white and alumina that were attracted to the left side, therefore positively charged. Figure 5 shows the results for the 4.71  $\mu$ m particles of alumina.

The particle charges in these cases were acquired by triboelectrification, i.e. the exchange of electrons during the contact of the particles with another solid (the duct walls or parts). In these cases, it is known that the exchange of electrons, hence the polarity, depends on the materials involved and their relative position in the triboelectric series (Hinds, 1999).

### **Charge Level**

#### **Mono-Dispersed Aerosol**

The results of the tests using the monodispersed aerosol generator are shown in Figure 6, where the



**Figure 4:** Total of pyrochlore particles as a function of position x, for  $d_{pS} = 1.93 \ \mu m$ .

particle charge is plotted as a function of the particle diameter. It can be noticed that the particle charge does not show any specific trend with the diameter. The complexity of the process of particle generation vibrating orifice, in which involves in the atomization of the solution, break-up of a jet to form evaporation of the droplets. droplets, and precipitation/crystallization/agglomeration of the solids, are certainly the cause of this high instability in the electrostatic charging.



Figure 5: Total of alumina particles as a function of position x, for  $d_{pS} = 4.71 \ \mu m$ .



Figure 6: Particle charge as a function of Stokes diameter (monodispersed aerosol).

### **Polydispersed Aerosols**

The results of the particle charge as a function of the diameter, using the rotating disc aerosol generator, are presented in Figures 7 to 12 for the polydispersed materials. The data show a linear dependence between particle diameter and the charge in these particles.

A linear relationship between charge and particle

diameter for the six powders analyzed can be expressed analytically by the parameters listed in Table 2. The linear correlation coefficients  $(r^2)$  obtained for all correlations are close to 0.99, indicating a good linearity for all the experimental data. These same linear behavior was earlier observed by Coury (1983) and Guang (1991), that worked with filtration of fly ash emitted by coal combustors.



Figure 7: Particle charge as function of Stokes diameter for concentrate phosphate rock.



Figure 9: charge as function of Stokes diameter for wood dust.



Figure 11: Particle charge as function of Stokes diameter for diatomite.



Figure 8: Particle charge as function of Stokes diameter for dry egg white.



Figure 10: Particle charge as function of Stokes diameter for alumina.



Figure 12: Particle charge as function of Stokes diameter for pyrochlore

Aerosol Charge (Q)  $r^2$  $-0.81 \times 10^{-11} d_{pS} + \overline{0.95 \times 10^{-1}}$ 0.986 Phosphate rock  $1.69 \times 10^{-11} d_{pS}^{r^2} - 2.28 \times 10^{-17}$ Alumina 0.993  $1.03 \times 10^{-11} d_{pS} - 3.01 \times 10^{-17}$ Dry egg white 0.950  $-1.88 \times 10^{-11} d_{pS}^{PS} + 2.17 \times 10^{-17}$ Wood dust 0.995  $\begin{array}{c} \text{-0.98} \times 10^{-11} d_{pS} + 0.37 \times 10^{-17} \\ \text{-0.62} \times 10^{-11} d_{pS} + 0.29 \times 10^{-17} \end{array}$ Pyrochlore 0.992 Diatomite 0.995

Table 2: Equations for particle charge as a functions of diameter

Figure 13 compares the level of charging (in absolute values) relative to particles with 4  $\mu$ m in diameter under the same experimental conditions. It can be said that the wood dust presents the highest level of charging (5,34x10<sup>-17</sup> C) while the dry egg white presents the lowest level of charge (1.11x10<sup>-17</sup> C) within the studied range.

Jonhston *et al.* (1987) proposed a power law expression for the absolute median particle charge with respect to particle diameter, in the form:

$$\left|\mathbf{Q}/\boldsymbol{e}\right| = \mathbf{A}\mathbf{d}_{\mathbf{p}}^{\mathbf{B}} \tag{4}$$

where e is the elementary charge of the electron, A is the median number of elementary charges of magnitude |e| on a particle of diameter 1 µm. Johnston found values of B typically in the range 1 to 2, and of A typically in the range 10 to 30. His results are given in Table 3, together with the results of the present work, adjusted to Equation 4. The amount of charge found in all cases was very similar, indicating their consistency. Note the methods of re-suspending dry powders in this Table involve mainly friction between the powder and the apparatus (triboelectrification) and consequently produced similar aerosols.

Table 3 also lists the experimental results obtained by Forsyth *et al.* (1998) for comparable aerosols obtained from liquids solutions. It can be seen that theirs results show a nearly linear dependence between particle charge and particle diameter. These trends, however, were not found in the present work for the methylene blue, also obtained from a solution.



Figure 13: Charge module value for particles 4.0 µm.

# Table 3: Power law fit, $|Q/e| = Ad_p^B$ , to generated aerosol charge distribution

Aerosol	Α	В	$\mathbf{R}^2$	Diameter range (um)	<b>Results from</b>			
From Dust Generator								
Phosphate rock (rotating disc)	48.6	1.06	0.97	0.15-8.0	this work			
Alumina (rotating disc)	7.83	2.58	0.85	1.4-4.7	this work			
Dry egg white (rotating disc)	1.07	2.74	0.98	2.8-8.5	this work			
Wood dust (rotating disc)	53.1	1.30	0.99	2.6-9.3	this work			
Pyrochlore (rotating disc)	29.79	1.48	0.96	0.9-5.4	this work			
Diatomite (rotating disc)	26.45	1.15	0.99	1.0-6.3	this work			
Coal (turntable-Venturi)	36.8	1.17	0.99	0.6-7.5	Johnston (1987)			
Quartz (turntable-Venturi)	27.8	1.48	0.99	0.6-7.5	Johnston (1987)			
Mica (turntable-Venturi)	30.3	1.18	0.99	0.6-7.5	Johnston (1987)			
From Liquid Solution								
Methilene blue (vibrating orifice)	N/A	N/A	N/A	6.0-14.0	this work			
NaCl (collision nebulizer)	4.6	1.32	0.92	0.6-2.1	Johnston (1987)			
KCl 0.1% (spray atomizer)	14.15	0.7	0.96	0.15-0.3	Forsyth (1998)			
KCl 1% (spray atomizer)	6.7	0.45	0.93	0.15-0.3	Forsyth (1998)			
NaCl 0.1%(spray atomizer)	19.1	0.8	0.96	0.15-0.75	Forsyth (1998)			
NaCl 10% (spray atomizer)	6.2	0.60	0.89	0.15-0.65	Forsyth (1998)			

#### CONCLUSIONS

The dependence of particle charge to particle diameter were measured for seven different materials. The values of the charge measured in the particles are of the same order of magnitude as the ones reported in the literature, indicating the adequacy of the equipment and of the method employed.

The mean Stokes diameters of the monodispersed particles studied range between  $6.0 \times 10^{-6}$  m and  $1.4 \times 10^{-5}$  m and the electric charges presented values from  $-3.0 \times 10^{-16}$  C to  $-5.0 \times 10^{-18}$  C. No evident dependence between the particle charge with its diameter was observed for the tests with the monodispersed aerosol, generated by a liquid solution.

For the tests using the poly-dispersed aerosols generated by triboelectrification, a linear dependence between particle charge and diameter was noticed. The analysis of the obtained data showed that the level of charging and polarity is a function of the powder composition: some aerosols were negatively charged (phosphate rock, diatomite, pyrochlore and wood dust) while others were positive (alumina and dry egg white). In the same experimental conditions, dust wood acquired five times more charge than egg white, in absolute values.

#### NOMENCLATURE

μ	Gas viscosity	kg.s-1.m-1
$\Delta V$	Electrical potential difference	V
$d_p$	Stokes diameter	m

Е	Electric field	V.m-1
Fd	Drag force	Ν
Fe	Electric force	Ν
Fs	Slip factor	(-)
L	Parallel plane plates distance	m
Q	Particle charge	С
$\mathrm{U}_0$	Gas velocity	m.s-1
U <sub>x</sub>	Particle velocity (direction <i>x</i> )	m.s-1
Х	Particle deflection	m
Ζ	Length of the deflection plates	m
Zp	Electromobility of the	$m^2.V^{-1}.s^{-1}$
•	particle	

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