

ENE 426 Final Report

Group D

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Introduction:

Air pollution is a serious challenge in Los Angeles, as it results in decreased visibility, degradation to buildings, and respiratory health problems. In this assignment, we sought to characterize common characteristics of particulate air pollution using data from the city.

This report includes particulate statistics, design of a cascade impactor to collect particulates of various sizes, an analysis of respiratory deposition, analysis of optical properties, analysis of condensational growth under a variety of relative humidity and salt conditions, and the design of an electrostatic precipitator (ESP) to control particulate emissions from an industrial plant.

These results of these steps will allow policy makers and officials to build pertinent policies to limit air pollution in LA. It will also investigate the efficiency of an ESP to limit these particulates. This project can also be applied to the current Coronavirus pandemic, as section C will illustrate how the virus can be spread and directly enter the lungs of a patient.

Part A: Particle Statistics (Sining Niu and Catherine Knox)

Introduction:

This particle size distribution data is retrieved from a real-world particle measurement in central Los Angeles. In part A, basic statistical analyses are conducted, and we could obtain a general interpretation of this data set.

Calculations and results:

First, particle number distribution could be calculated:

$$dN = \frac{dN}{d \log d_p} \times \Delta \log d_p$$

Then, total surface and volume and mass concentration can be obtained:

$$\text{Total surface concentration} = \sum(\pi \times d_p^2 \times dN) = \mathbf{5.96 \times 10^8 \text{ nm}^2/\text{cm}^3}$$

$$\text{Total volume concentration} = \sum\left(\frac{\pi}{6} \times d_p^3 \times dN\right) = \mathbf{2.87 \times 10^{10} \text{ nm}^3/\text{cm}^3}$$

$$\text{Total mass concentration} = \sum\left(\frac{\pi}{6} \times d_p^3 \times dN \times \rho\right) = \mathbf{2.88 \times 10^{-11} \text{ g/cm}^3}$$

Average and median diameters were obtained as following:

$$\text{Number based concentration average diameter} = \frac{\sum(N_i d_i)}{N} = \mathbf{71.82 \text{ nm}}$$

$$\text{Surface based concentration average diameter} = \left(\frac{\sum N_i d_i^2}{N}\right)^{1/2} = \mathbf{105 \text{ nm}}$$

$$\text{Volume/Mass based concentration average diameter} = \left(\frac{\sum N_i d_i^3}{N}\right)^{1/3} = \mathbf{150 \text{ nm}}$$

Calculation of median diameter based on geometric mean:

$$\text{Number based concentration median diameter} = \exp\left(\frac{\sum N_i \ln(d_i)}{N}\right) = \mathbf{47.9 \text{ nm}}$$

$$\text{Surface based concentration median diameter} = \exp\left(\frac{\sum S_i \ln(d_i)}{S}\right) = \mathbf{214.3 \text{ nm}}$$

$$\text{Volume/Mass based concentration median diameter} = \exp\left(\frac{\sum V_i \ln(d_i)}{V}\right) = \mathbf{418.4 \text{ nm}}$$

Calculation of median diameter with 50% cumulation:

$$\text{Number based concentration median diameter} = \mathbf{41.4 \text{ nm}}$$

$$\text{Surface based concentration median diameter} = \mathbf{229.7 \text{ nm}}$$

$$\text{Volume/Mass based concentration median diameter} = \mathbf{346 \text{ nm}}$$

Geometric standard deviation:

$$GSD = \frac{d_{50}}{d_{15.78}}$$

$$\text{Number based concentration GSD} = d_{50}/d_{15.78} = 41.40/19.50 = \mathbf{2.12}$$

$$\text{Surface based concentration GSD} = d_{50}/d_{15.78} = 229.70/98.20 = \mathbf{2.34}$$

$$\text{Volume/Mass based concentration GSD} = d_{50}/d_{15.78} = 346.00/192 = \mathbf{1.80}$$

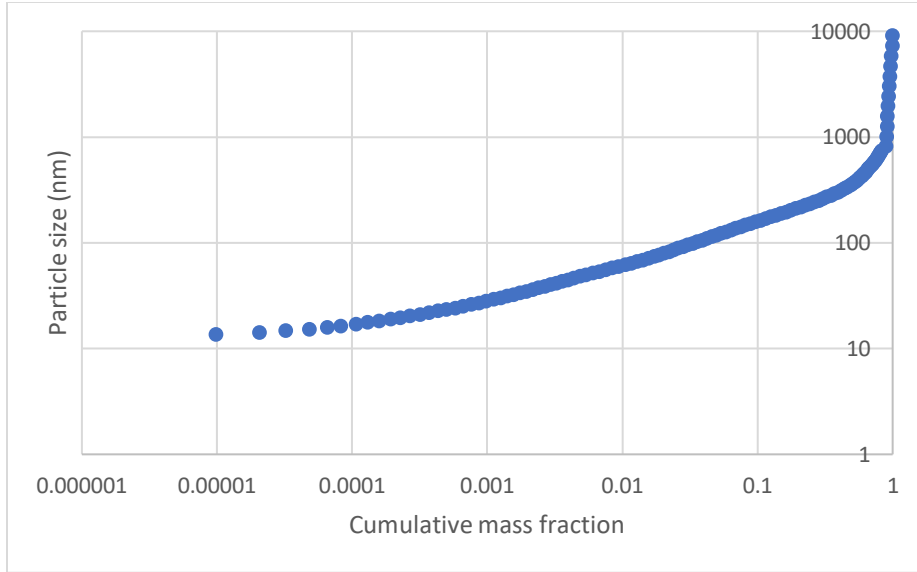


Figure A.1. Plot of particle size versus cumulative mass fraction in a log-log scale.

As we obtain the plot above, which is nearly a straight line, we can say that this data set is **lognormal distribution**.

Deposition:

$$t = 1 \text{ hr}, A = 1\text{m}^2, \lambda = 66.7\text{nm}$$

$$C_c = 1 + 1.7 \times 2 \times \frac{\lambda}{d_p}$$

Diffusion deposition:

$$D = \frac{kTC_c}{3\pi\eta d}$$

$$N(t) = 2n_0\left(\frac{Dt}{\pi}\right)^{0.5} A$$

Settling deposition:

$$V_{TS} = \frac{\rho_p d_p^2 g C_c}{18\mu}$$

$$N(t) = V_{TS} \times A \times t \times n_0$$

Calculation results are shown in the figures below:

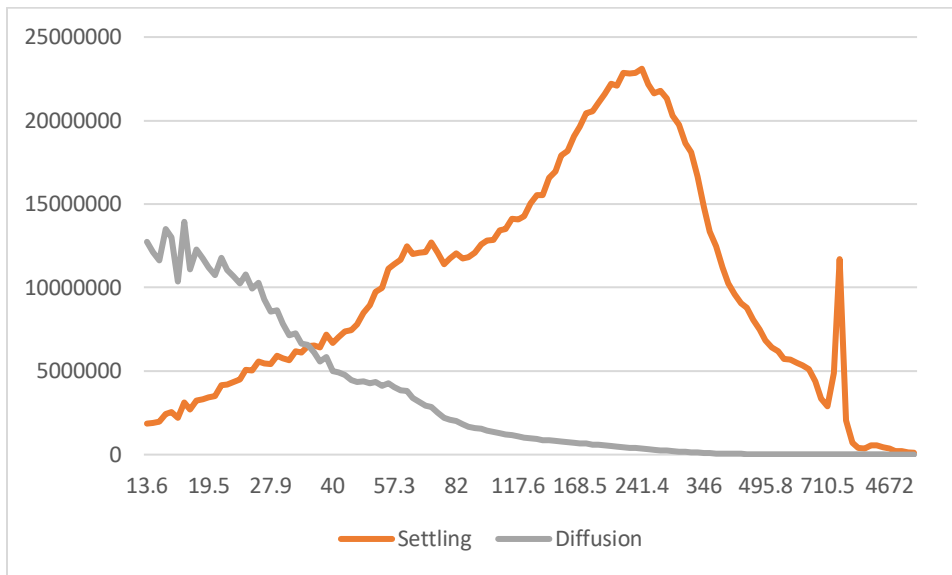


Figure A.2. Plot of settling and diffusion settling particle numbers at different particle size.

The outlier above for settling deposition around 800nm is due to a sudden increase of particle number concentration. This sudden increase might be the result of measure error.

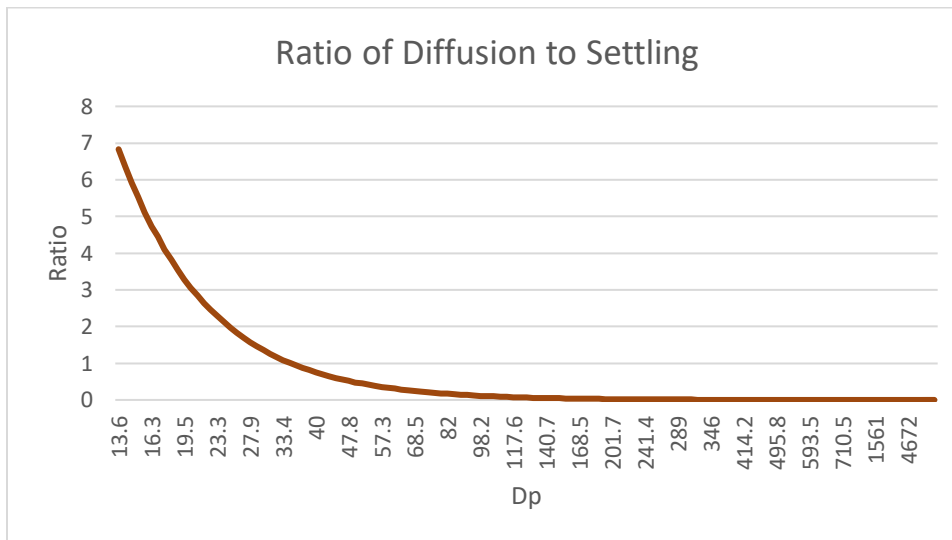


Figure A.3. Plot of the ratio of diffusion to settling deposition at a function of particle size.

As shown above in Figure A.3., the ratio of diffusion to settling deposition decreases as particle size increases. This is because the smaller particles have larger diffusivity and larger particles would prefer gravitational settling.

Part B: Cascade Impactor Design (Catherine Knox)

Introduction:

In this section, we will design a cascade impactor to capture 200 ug of particulates in each stage. This allows for further analysis of the Los Angeles particulates.

Calculations and Results:

Mass concentrations were obtained from Part A and shown in the table below:

Size	#Concentration/cm ³	Mass concentration (g/cm ³)
<0.2	16364	4.78E-12
0.2-0.5	1172	1.47E-11
0.5-1	48.5	6.39E-12
1-2.5	.89	9.66E-13
2.5-5	.039	9.54E-13
5-10	.005	7.42E-13

Table B.1. Mass concentrations for each selected bin size.

Based on these values, we design the impactor for the stage with the lowest mass concentration, given by 5-10 um at $7.42 \times 10^{-13} \text{ g/cm}^3$. Additionally, we know that the sampling period is 2 days and the required sample is 200 ug. Flow can be calculated as shown below:

$$\begin{aligned}
 \text{sample required} &= 200 \text{ ug} \\
 \text{sampling period} &= 2 \text{ days} \\
 M_{\min} &= QC_{\min}\Delta t \\
 C_{\min} \text{ occurs in stage 1: } &9.95 \times 10^{-7} \text{ ug/cm}^3 \\
 200 \text{ ug} &= Q (7.42 \times 10^{-7})(2 * 24 * 60) \\
 Q &= \frac{200 \text{ ug}}{\left(7.42 * \frac{10^{-7} \text{ ug}}{\text{cm}^3}\right) * 2880 \text{ minutes}} = 93590 \frac{\text{cm}^3}{\text{min}} = 1560 \frac{\text{cm}^3}{\text{s}} = .0936 \frac{\text{m}^3}{\text{min}} = 93.6 \text{ l/min}
 \end{aligned}$$

In order to calculate width, velocity, and pressure drop, we first have to assume length. The following equations were used to calculate these values:

$$\begin{aligned}
 Stk_{50} &= .59 \\
 d_{50}\sqrt{Cc} &= \left[\frac{9 * \eta * W^2 * L(Stk_{50})}{\rho p Q} \right]^{\frac{1}{2}} \\
 d_{50}\sqrt{Cc} &= d_{50} + .078
 \end{aligned}$$

$$\text{Pressure drop} = .5 * \rho * U^2$$

Given, this information, each stage was designed as shown:

Stage 1: 5-10 um

$$\begin{aligned}
 \text{Use cutpoint } D_p &= 5 \text{ um} \\
 C_c &= 1 + 1.7 * Kn/d_p
 \end{aligned}$$

$$Cc = 1 + 1.7 * (2 * 0.066)/5 = 1.045$$

$$d_{50}\sqrt{Cc} = \left[\frac{9 * \eta * W^2 * L(Stk50)}{\rho p Q} \right]^{\frac{1}{2}}$$

$$d_{50} = d_{50}\sqrt{Cc} - .078$$

$$d_{50} + .078 = \left[\frac{9 * \eta * W^2 * L(Stk50)}{\rho p Q} \right]^{\frac{1}{2}}$$

$$\text{Assume } L = 25 \text{ cm}$$

$$(5 + .078) * 10^{-4} = \left[\frac{9 * 1.8 * 10^{-4} * W^2 * 25 * 0.59}{1 * 1560} \right]^{1/2}$$

$$W = .13 \text{ cm}$$

$$U = \frac{Q_{max}}{W * L}$$

$$U = \frac{1560 \frac{cm^3}{s}}{25 \text{ cm} * .13 \text{ cm}} = 480 \text{ cm}^s$$

$$\text{Pressure drop} = .5 * \rho * U^2 = .5 * 1.2 * 10^{-3} (480^2) = 138.24 \frac{dyn}{cm^2}$$

Stage 2: 2.5-5 um

$$\text{Use cutpoint } D_p = 2.5 \text{ um}$$

$$Cc = 1 + 1.7 * Kn/dp$$

$$Cc = 1 + 1.7 * (2 * 0.066)/2.5 = 1.08976$$

$$d_{50} + .078 = \left[\frac{9 * \eta * W^2 * L(Stk50)}{\rho p Q} \right]^{\frac{1}{2}}$$

$$\text{Assume } L = 15 \text{ cm}$$

$$(2.5 + .078) * 10^{-4} = \left[\frac{9 * 1.8 * 10^{-4} * W^2 * 15 * 0.59}{1 * 1560} \right]^{1/2}$$

$$W = .085 \text{ cm}$$

$$U = \frac{Q_{max}}{W * L}$$

$$U = \frac{1560 \frac{cm^3}{s}}{15 \text{ cm} * .085 \text{ cm}} = 1223.5 \text{ cm}^s$$

$$\text{Pressure drop} = .5 * \rho * U^2 = .5 * 1.2 * 10^{-3} (1223.5^2) = 898.21 \frac{dyn}{cm^2}$$

Stage 3: 1-2.5 um

$$\text{Use cutpoint } D_p = 1 \text{ um}$$

$$Cc = 1 + 1.7 * Kn/dp$$

$$Cc = 1 + 1.7 * (2 * 0.066)/1) = 1.22$$

$$\text{Assume } L = 10 \text{ cm}$$

$$(1 + .078) * 10^{-4} = \left[\frac{9 * 1.8 * 10^{-4} * W^2 * 10 * 0.59}{1 * 1560} \right]^{1/2}$$

$$W = .04 \text{ cm}$$

$$U = \frac{Q_{max}}{W * L}$$

$$U = \frac{1560 \frac{cm^3}{s}}{10cm * .04 \text{ cm}} = 3900 \text{ cm}^s$$

$$\text{Pressure drop} = .5 * \rho * U^2 = .5 * 1.2 * 10^{-3} (3900^2) = 9126 \frac{dyn}{cm^2}$$

Stage 4: .5-1 um

$$\text{Use cutpoint } Dp = .5 \text{ um}$$

$$Cc = 1 + 1.7 * Kn/dp$$

$$Cc = 1 + 1.7 * (2 * 0.066)/.5) = 1.45$$

$$d_{50} + .078 = \left[\frac{9 * \eta * W^2 * L(Stk50)}{\rho p Q} \right]^{1/2}$$

$$\text{Assume } L = 10 \text{ cm}$$

$$(.5 + .078) * 10^{-4} = \left[\frac{9 * 1.8 * 10^{-4} * W^2 * 10 * 0.59}{1 * 1560} \right]^{1/2}$$

$$W = .023 \text{ cm}$$

$$U = \frac{Q_{max}}{W * L}$$

$$U = \frac{1560 \frac{cm^3}{s}}{10 \text{ cm} * .023 \text{ cm}} = 6782.6 \text{ cm}^s$$

$$\text{Pressure drop} = .5 * \rho * U^2 = .5 * 1.2 * 10^{-3} (6782.6^2) = 27,602 \frac{dyn}{cm^2}$$

Stage 5: .2-.5

$$\text{Use cutpoint } Dp = .2 \text{ um}$$

$$Cc = 1 + 1.7 * Kn/dp$$

$$Cc = 1 + 1.7 * (2 * 0.066)/.2) = 2.122$$

$$d_{50} + .078 = \left[\frac{9 * \eta * W^2 * L(Stk50)}{\rho p Q} \right]^{1/2}$$

$$\text{Assume } L = 15 \text{ cm}$$

$$(.2 + .078) * 10^{-4} = \left[\frac{9 * 1.8 * 10^{-4} * W^2 * 15 * 0.59}{1 * 1560} \right]^{1/2}$$

$$W = .01 \text{ cm}$$

$$U = \frac{Q_{max}}{W * L}$$

$$U = \frac{1560 \frac{\text{cm}^3}{\text{s}}}{15 \text{ cm} * .01 \text{ m}} = 10,400 \text{ cm}^s$$

$$\text{Pressure drop} = .5 * \rho * U^2 = .5 * 1.2 * 10^{-3} (10400^2) = 64,896 \frac{\text{dyn}}{\text{cm}^2}$$

Stage 6: <.2 um

Everything left in the flow

In order to verify that this system functions in terms of pressure drop, we must use the figures provided for the selected pump.

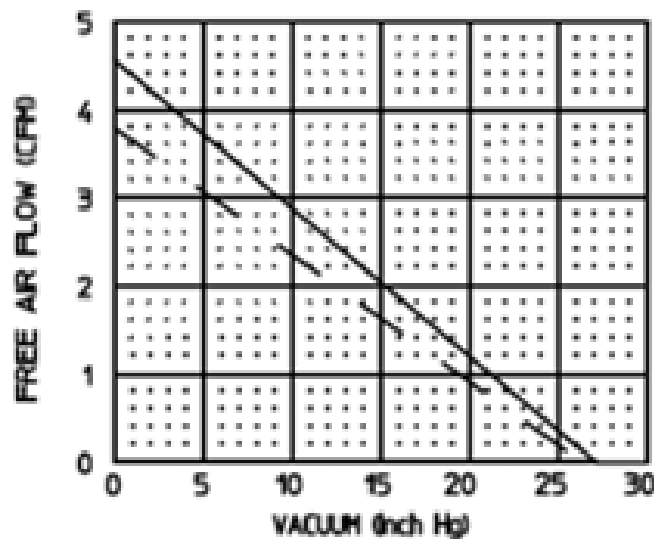


Figure B.1. Pressure chart for the selected pump

Given our flow rate of 3.31 CFM, we can see the maximum pressure is about 6 inches of mercury. Given that we operate the pump at around 85% of the maximum pressure, we can operate at 5.1 inches of mercury. Summing the values for pressure drop at each stage and adding the 10 inches of water drop from the filter results in .122 bar, or about 3.6 inches of mercury. Therefore, the design is feasible with all design parameters shown in the table below.

Stage	Length (cm)	Width (cm)	U (cm/s)	P (dyn/cm ²)
1 (5-10 um)	25	.13	480	138.20
2 (2.5-5 um)	15	.083	1,224	898.21
3 (1-2.5 um)	10	.04	3,100	9,126

4 (0.5-1 μm)	10	.023	6,783	27,602
5 (0.2-0.5 μm)	15	0.01	10,400	64,896
6 (<2)	Remaining flow	Remaining flow	Remaining Flow	Remaining Flow

Table B.2. Cascade impactor design parameters for each stage.

Part C: Respiratory System (Sining Niu)

Introduction:

Human respiratory system is divided into three different regions: head region (HA), tracheobronchial region (TB) and alveolar region (AL). Particles are deposited in respiratory system by interception, impaction and diffusion.

Calculations and results:

Respiratory deposition fraction calculations:

Head Region:

$$DF_{HA} = IF \left(\frac{1}{1 + \exp(6.84 + 1.183 \ln d_p)} + \frac{1}{1 + \exp(0.924 - 1.885 \ln d_p)} \right)$$
$$IF = 1 - 0.5 \left(1 - \frac{1}{1 + 0.00076 d_p^{2.8}} \right)$$

Tracheobronchial Region:

$$DF_{HA} = \left(\frac{0.00352}{d_p} \right) [\exp(-0.234(\ln d_p + 3.40)^2) + 63.9 \exp(-0.819(\ln d_p - 1.61)^2)]$$

Alveolar Region:

$$DF_{AL} = \left(\frac{0.0155}{d_p} \right) [\exp(-0.416(\ln d_p + 2.84)^2) + 19.411 \exp(-0.482(\ln d_p - 1.362)^2)]$$

Mass based deposition:

$$\text{mass based deposition} = DF \times \text{resp. rate} \times t \times \text{mass conc.}$$

Below in Table C.1, the final results for deposition fraction and mass-based deposition are shown.

Dp range (μm)	DF_HA	DF_TB	DF_AL	HA (μg)	TB (μg)	AL (μg)	Total (μg)
<0.2	0.021	0.027	0.142	2.10	2.63	14.09	18.82
0.2 – 0.5	0.056	0.005	0.064	16.93	1.43	19.38	37.74
0.5 – 1	0.189	0.016	0.109	26.44	2.27	15.31	44.02
1 – 2.5	0.532	0.052	0.126	12.95	1.27	3.07	17.29
2.5 – 5	0.815	0.056	0.080	19.58	1.35	1.92	22.85
5 – 10	0.863	0.026	0.033	16.33	0.50	0.62	17.45

Table C.1. Results for deposition fraction and mass-based deposition.

From the results above, we can see that larger particles are more likely to deposited in head region by impaction. Since the air flow speed is relatively high there. While for smaller and ultrafine particles, they can reach the tracheobronchial and alveolar region due to diffusion, since the air flow is calmer in lower lung. This coronavirus could adhere to small particles and then easily penetrate into our respiratory system by diffusion.

Recalculated with RH = 90%, salt mass fraction = 25% & 75%

First, need to recalculate the mass concentration for each stage with the new diameter (Dp, retrieved from part E) and origin particle number concentration (Ni):

$$Mass\ Conc. = Ni \times \frac{\pi}{6} \times d_p^3 \times \rho_p$$

Then, recalculate deposition fraction and mass-based deposition with the same method and results are shown below:

Respiratory Deposition RH = 90%, Salt = 25%								
Midpoint Dp (μm)	Mass conc. (μg/m ³)	DF_HA	DF_TB	DF_AL	HA (μg)	TB (μg)	AL (μg)	Total (μg)
0.100	8.52	0.021	0.026	0.141	3.61	4.50	24.09	32.20
0.352	26.18	0.056	0.005	0.064	29.36	2.47	33.47	65.30
0.753	10.52	0.190	0.016	0.110	40.04	3.44	23.09	66.57
1.758	2.26	0.534	0.052	0.126	24.10	2.36	5.68	32.14
3.766	0.97	0.816	0.056	0.080	15.91	1.09	1.56	18.56
7.532	0.92	0.863	0.026	0.032	15.86	0.48	0.60	16.94

Table C.2. Results for deposition fraction and mass-based deposition with RH = 90% and salt mass fraction = 25%.

Respiratory Deposition RH = 90%, Salt = 75%								
Midpoint Dp (μm)	Mass conc. (μg/m ³)	DF_HA	DF_TB	DF_AL	HA (μg)	TB (μg)	AL (μg)	Total (μg)
0.145	8.52	0.021	0.015	0.087	10.59	7.53	44.52	62.64
0.507	26.18	0.102	0.007	0.084	159.76	11.46	131.96	303.18
1.086	10.52	0.318	0.031	0.126	200.66	19.48	79.74	299.88
2.535	2.26	0.693	0.061	0.109	93.77	8.23	14.68	116.68
5.432	0.97	0.870	0.041	0.053	50.87	2.41	3.07	56.35
10.864	0.92	0.789	0.013	0.017	43.53	0.70	0.92	45.15

Table C.3. Results for deposition fraction and mass-based deposition with RH = 90% and salt mass fraction = 75%.

Similar conclusion can be obtained that larger particle are more likely to deposit in head region. Besides, as particle size increases, from mass-based perspective, much more particle mass would deposit in our lung.

Part D: Optical Properties (Marco Kleimans)

Introduction:

Driven by the scattering and the absorption of light, optical properties of particles are in charge of many atmospheric phenomena, such as rainbow and smog formation, and visibility degradation, among others. The latter is the focus of this part.

Calculations and results:

In order to calculate visibility, we need to know the **extinction coefficient (Q_e)**, which is a coefficient attributed to the attenuation of light due to the combined effects of **absorption** and **scattering (σ_e)**. Since absorption is ignored, we thus have:

$$Q_e = \sigma_e$$

We will use as an example on how the calculations were performed with the unaffected particles midpoint of the first stage (0 – 200 nm)¹. This value corresponds to a median diameter of 100 nm. All the remaining values will be obtained following this method and the results will be reported in tables/graphs below:

First, we need to find the *size parameter* (α):

$$\alpha = \frac{\pi * d}{\lambda} \quad (\text{where } \lambda \text{ is the mid value of visible light, or 550 nm})$$

So,

$$\alpha = \frac{\pi * 100}{550} = \mathbf{0.571}$$

Then, we compute *Q_e* (the *extinction coefficient*) from the provided website in the description of the project:

$$Q_e \text{ (from website): } \mathbf{0.025^2}$$

It is important to note that for particles where $d_p > 4000$ nm, the *Q_e* value converges to **2** because of the extinction efficiency paradox.

Next, we calculate the scattering coefficient (σ_e):

$$\sigma_e = \sum_i \frac{\pi N_i d_i^2 (Q_e)_i}{4}$$

So,

$$\sigma_e = \frac{(16364 \frac{\text{particles}}{\text{cm}^3}) \left(\frac{100 \text{ nm}}{1 \text{ m}}\right)^3 (100 \text{ nm})^2 \left(\frac{1 \text{ m}}{10^9 \text{ nm}}\right)^2 (0.025)}{4}$$

$$\sigma_e = \mathbf{3.18 \times 10^6 m^{-1}}$$

The final step is to determine *visibility* (L_v):

¹ Results for unaffected particles are not reported because they are irrelevant in the analysis. However, the data is available in the attached Excel file (i.e. Excel Part_D_Group_D)

² Calculations obtained from http://irina.eas.gatech.edu/Lab_Source/sphere5235.aspx

$$L_v = \frac{-\ln(0.02)}{\sigma_e} = \frac{3.91}{\sigma_e}$$

So,

$$L_v = \frac{3.91}{3.18 \times 10^6 m^{-1} \left(\frac{1000m}{1km} \right)}$$

$$L_v = 1228.083419 \text{ km}$$

Finally, we calculate overall visibility. For this part, the σ_e values for each individual midpoint value for each cut-off range have been summed up using the sum function in excel (i.e =sum(σ_{0-200} : $\sigma_{5000-10000}$)) and then the summed value was evaluate using the formula in step 4. For instance:

Midpoint (dp [=] nm)	Scattering coefficient (σ [=] m ⁻¹)	Visibility (Lv [=] km)
100	3.18E-06	1228.083419
350	2.03E-04	19.2996188
750	9.32E-05	41.96975745
1750	6.19E-06	631.8590341
3750	9.41E-07	4152.972698
7500	4.42E-07	8854.483129
Summed σ		3.07E-04
Overall Visibility		12.75643426

Table D.1. Summed values for the scattering of light and summation across scattering coefficient column to obtain overall visibility (unaffected particles).

These are the results obtain:

Optical Properties - Salt 25%							
	Stage (dp [=] nm)	Midpoint (dp [=] nm)	Ni (particles/cm ³)	α	QE	Scattering coefficient (σ [=] m ⁻¹)	Visibility (Lv [=] km)
RH 95%	0 - 200	128.837	16364.15385	0.736	0.068	1.44E-05	270.68
	200 - 500	450.931	1172.06938	2.576	2.729	5.11E-04	7.66
	500 - 1000	966.281	48.47386	5.519	3.155	1.12E-04	34.86
	1000-2500	2254.655	8.92E-01	12.879	2.179	7.76E-06	504.05
	2500 - 5000	4831.404	3.91E-02	27.597	2.000	1.43E-06	2726.41
	5000 - 10000	9662.809	5.00E-03	55.194	2.000	7.33E-07	5334.32
RH 90%	0 - 200	100.4319169	16364.15385	0.574	0.025	3.28E-06	1192.34
	200 - 500	351.5117093	1172.06938	2.008	1.813	2.06E-04	18.96
	500 - 1000	753.2393771	48.47386	4.302	4.360	9.42E-05	41.52
	1000-2500	1757.558547	8.92E-01	10.039	2.840	6.14E-06	636.48
	2500 - 5000	3766.196886	3.91E-02	21.512	2.133	9.29E-07	4206.76
	5000 - 10000	7532.393771	5.00E-03	43.025	2.000	4.45E-07	8778.49
RH 80%	0 - 200	100	16364.15385	0.571	0.025	3.18E-06	1228.08
	200 - 500	268.254	1172.06938	1.532	0.801	5.31E-05	73.70
	500 - 1000	574.829	48.47386	3.283	3.644	4.58E-05	85.30
	1000-2500	1341.268	8.92E-01	7.661	1.796	2.26E-06	1728.28
	2500 - 5000	2874.146	3.91E-02	16.417	2.499	6.34E-07	6165.36
	5000 - 10000	5748.292	5.00E-03	32.834	2.000	2.59E-07	15073.29

Table D.2.a. Results for all the parameters for Salt 25%.

Optical Properties - Salt 75%							
	Stage (dp [=] nm)	Midpoint (dp [=] nm)	Ni (particles/cm ³)	α	QE	Scattering coefficient (σ [=] m ⁻¹)	Visibility (Lv [=] km)
RH 95%	0 - 200	185.815755	16364.15385	1.061	0.264	1.17E-04	33.32050816
	200 - 500	650.3551424	1172.06938	3.715	4.178	1.63E-03	2.40381461
	500 - 1000	1393.618162	48.47386	7.960	1.691	1.25E-04	31.27352634
	1000-2500	3251.775712	8.92E-01	18.574	2.366	1.75E-05	223.218674
	2500 - 5000	6968.090812	3.91E-02	39.802	2.000	2.98E-06	1310.720599
	5000 - 10000	13936.18162	5.00E-03	79.603	2.000	1.52E-06	2564.47309
RH 90%	0 - 200	144.8478891	16364.15385	0.827	0.106	2.87E-05	136.31
	200 - 500	506.9676117	1172.06938	2.896	3.362	7.95E-04	4.92
	500 - 1000	1086.359168	48.47386	6.205	2.477	1.11E-04	35.13
	1000-2500	2534.838059	8.92E-01	14.479	2.068	9.31E-06	420.17
	2500 - 5000	5431.79584	3.91E-02	31.026	2.000	1.81E-06	2157.00
	5000 - 10000	10863.59168	5.00E-03	62.053	2.000	9.26E-07	4220.26
RH 80%	0 - 200	110.5396	16364.15385	0.631	0.037	5.79E-06	675.08
	200 - 500	386.8887	1172.06938	2.210	2.087	2.88E-04	13.60
	500 - 1000	829.0472	48.47386	4.736	3.861	1.01E-04	38.70
	1000-2500	1934.4435	8.92E-01	11.050	3.014	7.90E-06	495.10
	2500 - 5000	4145.236	3.91E-02	23.678	2.000	1.06E-06	3703.73
	5000 - 10000	8290.4721	5.00E-03	47.355	2.000	5.40E-07	7246.48

Table D.2.b. Results for all the parameters for Salt 75%.

Overall Visibility		
Relative Humidity	Salt 25%	Salt 75%
0.95	6.040446481	2.067714644
0.9	12.56519675	4.127158916
0.8	37.15666169	9.682361663

Table D.3. Overall visibility of both salt types across the different relative humidity values.

From these data, the following trends can be observed. First and foremost, as it can be appreciated from table D.3, that as relative humidity increases, the visibility decreases, and particles which are 75% salt have a greater impact on overall visibility. This trend is illustrated in the following graph:

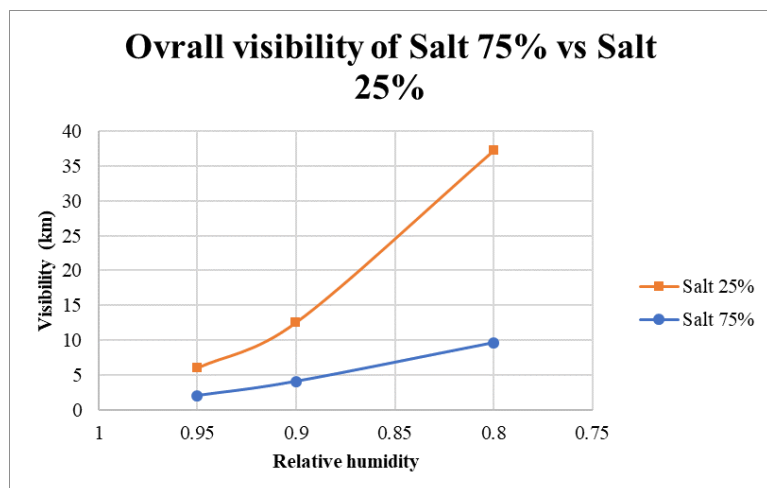


Figure D.1. As it can be seen from this plot, the visibility is affected by both relative humidity and salt percent of each type of particle.

Furthermore, visibility reaches a minimum at the accumulation mode particles. When observing table D.2.a and D.2.b, one can see that the trend is consistent across salt types and humidity conditions. This is

because particles of this size do not have effective mechanisms to deposit (i.e. these are too massive for diffusion and too small for gravitational settling) and therefore remain suspended in the air for longer periods of time. This trend can be graphically appreciated below.

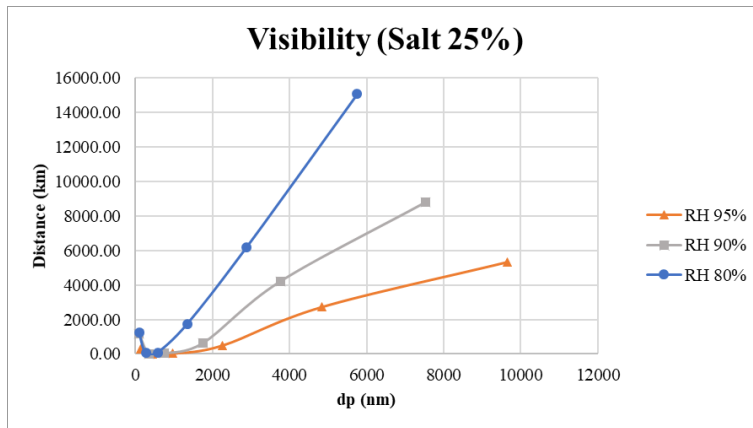


Figure D.2.a. Visibility of salt 25% at the different stages in the different relative humidity conditions.

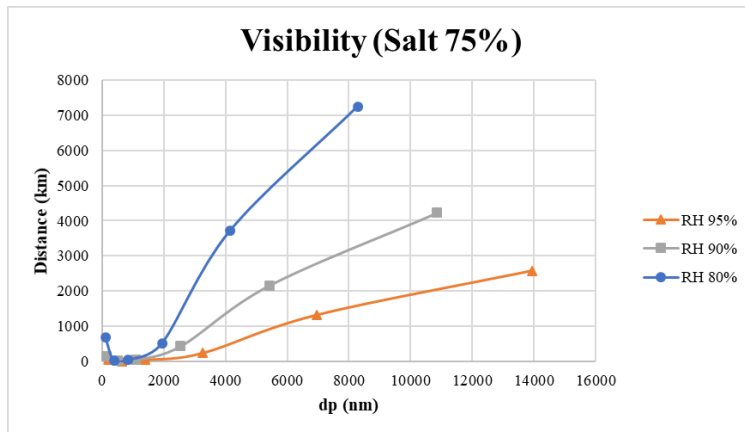


Figure D.2.b. Visibility of salt 75% at the different stages in the different relative humidity conditions.

Part E: Condensational growth (Zijiao Zhang)

Introduction:

In this part, we will assume that each size range has hygroscopic components that are 25% and 75% by mass. Then we need to calculate growth in lungs assuming 95% relative humidity in our respiratory tract and redo that for relative humidity equal to 80 and 90%. We also need to calculate surface area and mass or volume concentrations of grown aerosol. For the parts of respiratory deposition, scattering calculations and visibility range calculations, we have calculated the values in the previous parts.

Calculations and results:

For the first step, we need to calculate the grown particle diameter. Based on Kohler equation, we can ignore Kelvin effect and just consider the Raoult term. And the equation is posted below.

$$\frac{p}{P_s} = \left(1 + \frac{6im_sM_w}{M_s\rho_w\pi d_{(p,grown)}^3} \right)^{-1}$$

We already know the number of dissolved ions or i is equal to 2. Molecular weight of water or M_w is equal to 18. Molecular weight of ammonium nitrate or M_s is equal to 80. And the density of water or ρ_w is equal to 1 g/cm^3 . So we can substitute these given data in different conditions where the salt mass fraction are 25 percent and 75%, and when relative humidity are 95%, 90% and 80%. And then the grown particle diameter or d_p can be calculated.

For example, when the salt percentage is 25% and the relative humidity is 95%, we can get this equation below and calculate grown particle diameter.

$$\frac{p}{P_s} = 95\% = \left(1 + \frac{6im_sM_w}{M_s\rho_w\pi d_{(p,grown)}^3} \right)^{-1} = \left(1 + \frac{6 \times 2 \times 25\% \times 1.72 \times \frac{\pi}{6} d_p^3 \times 18}{80 \times 1 \times \pi \times d_{(p,grown)}^3} \right)^{-1}$$

So these are results when the salt mass fraction is 25% as table E.1.(a). shows. We can see the grown particle diameters for different relative humidity.

d_p (μm)	m_s (g)	$d_{(p,grown)}$, 80% RH (μm)	$d_{(p,grown)}$, 90% RH (μm)	$d_{(p,grown)}$, 95% RH (μm)
0.1	1.309E-16	0.08	0.10	0.13
0.35	5.61232E-15	0.27	0.35	0.45
0.75	5.52233E-14	0.57	0.75	0.97
1.75	7.01541E-13	1.34	1.76	2.25
3.75	6.90291E-12	2.87	3.77	4.83
7.5	5.52233E-11	5.75	7.53	9.66

Table E.1.(a). Results of grown diameter for salt 25%.

And these are results when the condition is 75% hygroscopic components for different relative humidity as table E.1.(b). shows.

d_p (μm)	m_s (g)	$d_{(p,\text{grown})}$, 80% RH (μm)	$d_{(p,\text{grown})}$, 90% RH (μm)	$d_{(p,\text{grown})}$, 95% RH (μm)
0.1	3.92699E-16	0.11	0.14	0.19
0.35	1.6837E-14	0.39	0.51	0.65
0.75	1.6567E-13	0.83	1.09	1.39
1.75	2.10462E-12	1.93	2.53	3.25
3.75	2.07087E-11	4.15	5.43	6.97
7.5	1.6567E-10	8.29	10.86	13.94

Table E.1.(b). Results of grown diameter for salt 75%.

Compare with these two charts, we can conclude that in the same salt percentage, the grown particle diameter also increases with the relative humidity increasing. And in the same relative humidity percentage, with the salt percentage increasing, grown particle diameter is increasing too.

For part of mass or volume concentrations and surface area of grown aerosol calculations, the number concentration of the particles stays the same as part A. We already know the equations of volume concentrations and surface area as follows.

For every particle size:

$$dN = \frac{dN}{d \log d_p} \times (\log D_{p,u} - \log D_{p,l})$$

$$\text{Mass concentration} = dN \times \frac{\pi}{6} \times d_p^3 \times \rho_p$$

$$\text{Volume concentration} = dN \times \frac{\pi}{6} \times d_p^3$$

$$\text{Surface area} = \pi \times d_p^2 \times dN$$

Total mass concentration, volume concentration and surface area are given by following equations:

$$\text{Total mass concentration} = \sum \text{Mass concentration of every particle size}$$

$$\text{Total volume concentration} = \sum \text{Volume concentration of every particle size}$$

$$\text{Total surface area} = \sum \text{Surface area of every particle size}$$

So we can get the values for 25% hygroscopic components and for different relative humidity which are 80% and 90% as table E.2.(a). shows.

	RH = 80%			RH = 90%		
d_p (μm)	d_(p,grown) (μm)	Volume Concentration (μm³/m³)	Surface Area (μm²/m³)	d_(p,grown) (μm)	Volume Concentration (μm³/m³)	Surface Area (μm²/m³)
0.1	0.08	3.77E+06	2.95E+08	0.10	8.49E+06	5.07E+08
0.35	0.27	1.12E+07	2.50E+08	0.35	2.52E+07	4.29E+08
0.75	0.57	4.18E+06	4.36E+07	0.75	9.40E+06	7.49E+07
1.75	1.34	6.63E+05	2.97E+06	1.76	1.49E+06	5.09E+06
3.75	2.87	3.26E+05	6.80E+05	3.77	7.33E+05	1.17E+06
7.5	5.75	3.50E+05	3.65E+05	7.53	7.88E+05	6.27E+05

Table E.2.(a). Results for 25% hygroscopic components and for different relative humidity.

And next chart is the results for 75% hygroscopic components and for different relative humidity which are 80% and 90% as table E.2.(b). shows.

	RH = 80%			RH = 90%		
d_p (μm)	d_(p,grown) (μm)	Volume Concentration (μm³/m³)	Surface Area (μm²/m³)	d_(p,grown) (μm)	Volume Concentration (μm³/m³)	Surface Area (μm²/m³)
0.1	0.08	3.77E+06	2.95E+08	0.10	8.49E+06	5.07E+08
0.35	0.27	1.12E+07	2.50E+08	0.35	2.52E+07	4.29E+08
0.75	0.57	4.18E+06	4.36E+07	0.75	9.40E+06	7.49E+07
1.75	1.34	6.63E+05	2.97E+06	1.76	1.49E+06	5.09E+06
3.75	2.87	3.26E+05	6.80E+05	3.77	7.33E+05	1.17E+06
7.5	5.75	3.50E+05	3.65E+05	7.53	7.88E+05	6.27E+05

Table E.2.(b). Results for 75% hygroscopic components and for different relative humidity.

Part F: Electrostatic Precipitator (Marco Kleimans)

Introduction:

Electrostatic precipitators take advantage of the electrical properties of aerosols to concentrate particles of desired sizes. The objective of this part is to analyze the parameters for the design of an electrostatic precipitator.

Calculations and results:

In order to begin the calculations, the following assumptions were made:

1. Electric field [V/m] = 60,000
2. Dielectric constant (ϵ) = 5
3. Minimum collection efficiency = 85%
4. Plate Height [m] = 5
5. Plate Width [m] = 6
6. Distance between plates [m] = 0.2
7. Flow rate [m^3/s] = 250
8. Energy cost [\$/kWh] = 0.12

These were data given in the problem statement. There are also other fundamental variables that need to be considered. These are the following:

Variables	
<i>k-constant (ft)</i>	0.6
<i>Ion mobility (m^2/V)</i>	1.50E-04
<i>Charge on an electron (C)</i>	1.60E-19
<i>Mean Thermal Speed of Ions (m/s)</i>	240
<i>Proportionality Constant K_e ($\text{N}\cdot\text{m}^2/\text{C}^2$)</i>	9.00E+09
<i>$Ni\cdot t$ (ions/m^3)</i>	1E+13
<i>Boltzmann constant ($\text{m}^2\cdot\text{kg}\cdot\text{s}^{-2}\cdot\text{K}^{-1}$)</i>	1.38E-23
<i>Temperature (K)</i>	293
<i>Mean Free Path</i>	0.066
<i>Viscosity of air ($\text{kg}/(\text{m}\cdot\text{s})$)</i>	1.80E-05
<i>Surface area total (m)</i>	60

Table F.1. Important constants required for calculations.

The following steps were followed for the design of the ESP. First, we computed the power to reach 85% efficiency:

$$E = 1 - \exp\left(-\frac{kPc}{Q}\right) \Rightarrow Pc = \frac{-Q}{k} \ln(1 - E)$$

So,

$$Pc = -\frac{8827 \frac{\text{ft}^3}{\text{s}}}{0.6} \ln(1 - 0.85)$$

$$Pc = 27910 \text{ W}$$

Then, we determined the potential (dW [V]) and the current (I [A]) in the ESP:

$$a. dW = \vec{E} * dx = (60,000)(0.2) = \mathbf{12,000V}$$

$$b. I = \frac{Pc}{dW} = \frac{27,910}{12,000} = \mathbf{2.3A}$$

At this point, we will use as an example on how the calculations were performed with the midpoint of the unaffected particle at the first stage (0 – 200 nm)³, similar to part D. This value corresponds to a median diameter of 100 nm. All of the remaining values will follow this method and the results will be reported in tables/graphs below. The following calculation methods are applicable for all size ranges, across the different types of salt and humidity conditions.

First, to calculate the diffusion and the field charging, we assumed an initial concentration (N_i [#*s*cm⁻³]) of 10¹³. Total charging was obtained from the addition of field charging $n_f(t)$ and diffusion charging $n_d(t)$ as follows:

$$n(t) = \frac{d_p kT}{2K_E e^2} \ln \left[1 + \frac{\pi K_E d_p \bar{c}_i e^2 N_i t}{2kT} \right]$$

So,

$$n_d(t) = \frac{(100 \text{ nm}) \left(\frac{10^{-9} \text{ m}}{1 \text{ nm}} \right) (1.38 \times 10^{-23}) (293 \text{ K})}{2(9 \times 10^9)(1.6 \times 10^{-19})^2} \ln \left[1 + \frac{\pi (9 \times 10^9)(100 \text{ nm}) \left(\frac{10^{-9} \text{ m}}{1 \text{ nm}} \right) (240)(1.6 \times 10^{-19})^2 (\mathbf{10^{13}})}{2(1.38 \times 10^{-23})(293 \text{ K})} \right]$$

$$n_d(t) = \mathbf{2.73}$$

And,

$$n(t) = \left(\frac{3\varepsilon}{\varepsilon + 2} \right) \left(\frac{Ed_p^2}{4K_E e} \right) \left(\frac{\pi K_E e Z_i N_{i\infty} t}{1 + \pi K_E e Z_i N_{i\infty} t} \right)$$

So,

$$n_f(t) = \left(\frac{3 * 5}{5 + 2} \right) \left(\frac{(100 \text{ nm})^2 \left(\frac{10^{-9} \text{ m}}{1 \text{ nm}} \right)^2 (60,000)}{4(9 \times 10^9)(1.6 \times 10^{-19})} \right) \left(\frac{\pi (9 \times 10^9)(1.6 \times 10^{-19})(1.5 \times 10^{-4})(\mathbf{10^{13}})}{1 + \pi (9 \times 10^9)(1.6 \times 10^{-19})(1.5 \times 10^{-4})(\mathbf{10^{13}})} \right)$$

$$n_f(t) = \mathbf{0.19}$$

Thus, the total charge concentration is:

$$n(t) = n_d(t) + n_f(t) = 2.73 + 0.19$$

$$\mathbf{n(t) = 2.93}$$

³Results for unaffected particles are not reported because they are irrelevant in the analysis. However, the data is available in the attached Excel file (i.e. Excel Part_F_Group_D)

Next, we needed to calculate the total area of the plates (A_p) in order to have the targeted collection efficiency of 85%. To do so, we were required to determine the following first:

a. Charge (q):

$$q = e * n(t) = (1.6 \times 10^{-19}) * 2.93 = 4.68 \times 10^{-19} C$$

b. Slip Correction Factor (C_c):

$$C_c = 1 + \frac{2.52\lambda}{d} = 1 + \frac{2.52 (0.066)}{100 \text{ nm} \left(\frac{10^{-3} \mu m}{1 \text{ nm}} \right)}$$

$$C_c = \mathbf{2.66}$$

c. Particle mobility (B)

$$B = \frac{C_c}{3\pi\eta d} = \frac{2.66}{3\pi(1.8 \times 10^{-5})(100 \text{ nm}) \left(\frac{10^{-9} \text{ m}}{1 \text{ nm}} \right)}$$

$$\mathbf{B = 1.57 \times 10^{11} \frac{N}{m \cdot s}}$$

d. Electric mobility (Z)

$$Z = qB = (4.68 \times 10^{-19})(1.57 \times 10^{11})$$

$$\mathbf{Z = 7.35 \times 10^{-8} \frac{m^2}{V \cdot s}}$$

e. Settling velocity (V_{te})

$$V_{TE} = ZE = 7.35 \times 10^{-8} * 60,000$$

$$\mathbf{V_{TE} = 0.0044 \frac{m}{s}}$$

It is important to note that many of these calculations can be replaced by data in Appendix A.11 from Hinds, but since the diameter sizes are specific, these cannot be located on the table. Therefore, for the sake of consistency, all the values were derived from calculations.

Once settling velocity was obtained, the preferred total plate area was calculated:

$$E = 1 - \exp\left(-\frac{V_{TE} A_c}{Q}\right) \Rightarrow A_p = -\frac{Q}{V_{TE}} \ln(1 - E)$$

$$A_p = -\frac{250}{0.0044} \ln(1 - 0.85)$$

$$\mathbf{A_p = 107527 \text{ m}^2}$$

After having found the areas for each of the range sizes, the largest value was selected, which corresponded to **126689 m²** for the 200-500nm stage. This can be observed in the following table.

Stage (dp [=] nm)	Settling Velocity (V/m)	Ac (m)
100 - 200	0.0044	107527.2242
200 - 500	0.0037	126689.0246
500 - 1000	0.0041	115767.8007
1000-2500	0.0053	88875.79008
2500 - 5000	0.0078	60933.65102
5000 - 10000	0.0122	38907.75067

Table F.2. Different plate areas for each stage given the corresponding settling velocity (unaffected particles). The largest area (200-500 nm) was selected.

Now, we computed the new ion concentration:

$$N_i = \frac{I}{A_p * E * Z_i * e}$$

So,

$$N_i = \frac{2.3}{(126689)(250) * (1.5 \times 10^{-4}) * (1.5 \times 10^{-19})}$$

$$N_i = 1.27 \times 10^{13}$$

We also found the number of plates:

$$N_p = \frac{A_p}{2 * \text{Area of single plate}} = \frac{126689}{30 * 2}$$

$$N_p = 2112$$

Then, we evaluated the airflow velocity:

$$U = Q / (\text{number of plates} - 1) / (H * D_x)$$

So,

$$U = 250 / (2112 - 1) / (5 * 0.2)$$

$$U = 0.118 \text{ m/s}$$

Next, we calculated the residence time:

$$\tau = L / U = 6 / 0.118$$

$$\tau = 50.6 \text{ s}$$

Once we obtained the new particle concentration (N_i) and the residence time (τ), we were able to calculate the new particle concentration N_{it} :

$$N_{it} = N_i * \tau = 1.27 \times 10^{13} * 50.6$$

$$N_{it} (\text{new}) = 6.5 \times 10^{14}$$

When the new N_{it} is obtained, the calculations are repeated, from the evaluation of the total charges to the computation of – once again – a second N_{it} . When calculating the plate area, we updated to the new efficiency (97%) and the most-recently calculated terminal velocity (V_{te}), corresponding to the previously

calculated plate area. This way we can confirm that our updated efficiency is correct (the Nit from both steps must match):

$$\text{Nit (check)} = 6.5 \times 10^{14}$$

At this point, as stated before, we also calculated the updated efficiency:

$$E = 1 - \exp\left(-\frac{V_{TE} A_c}{Q}\right)$$

where V_{te} comes from the latest iteration with the latest Nit value and the plate area (A_p) corresponds to the value from the first iteration.

$$E = 1 - \exp\left(-\frac{(0.069)(126689)}{(250)}\right)$$

$$E = 97\%$$

Finally, we estimated the purchasing and cost with the updated particle concentration and plate area:

$$\text{Purchasing cost} = a * A_p^b \text{ where } a = 96.0 \text{ and } b = 0.63 \text{ and } A_p \text{ in } ft^2$$

So,

$$\text{Purchasing cost} = 96 * \left(128598 m^2 * 10.7639 \frac{ft^2}{m^2}\right)^2$$

$$\text{Purchasing cost} = \$709,953.61$$

Similarly, we can calculate the operational costs:

$$\text{Operational cost} = Pc * \frac{\$}{kWh} * hr$$

So,

$$\text{Daily operational cost} = 27910 W \left(\frac{1 kW}{1000 W}\right) \left(\frac{\$0.12}{1 kWh}\right) (24hr)$$

$$\text{Daily operational cost (DOC)} = \$80.40$$

And,

$$\text{Yearly operational cost} = \text{DOC} * 365$$

So,

$$\text{Yearly operational cost} = \$80.40 * 365$$

$$\text{Yearly operational cost (YOC)} = \$29,344.32$$

These are the results obtained for every particle type at different relative humidity conditions:

		SALT 25%												
		Stage (dp [=] nm)	Midpoint (dp [=] nm)	Midpoint (dp [=] m)	Diffusion Charging	Field Charging	Total Charges	Charge (C)	Particle Mobility (m ² /Vs)	Slip Correction	Electric Mobility (m ² /Vs)	Settling Velocity (V/m)	Ac (m)	
RH - 95%	First iteration	100 - 200	128.837	1.29E-07		3.80	0.32	4.12	6.58923E-19	1.04816E+11	2.290933505	6.90658E-08	0.004143947	114451.27
		200 - 500	450.931	4.51E-07		18.14	3.96	22.10	3.53584E-18	1.36836917	6.32689E-08	0.003796134	124937.65	
		500 - 1000	966.281	9.66E-07		45.30	18.16	63.46	1.01538E-17	718035637	1.17212844	7.26034E-08	0.00456204	108874.60
		1000-2500	2254.655	2.25E-06		122.41	98.90	221.31	3.54089E-17	2807284827	1.073767383	9.94028E-08	0.00596417	79521.54
		2500 - 5000	4831.404	4.83E-06		294.58	454.12	748.70	1.19792E-16	1262065593	1.03424776	1.51185E-07	0.009071127	52284.57
	5000 - 10000	9662.809	9.66E-06		647.93	1816.46	2464.39	3.94303E-16	620532614	1.017212386	2.44678E-07	0.014680657	32306.46	
	Second iteration	100 - 200	128.837	1.29E-07		8.47	0.37	8.84	1.41E-18	1.05E+11	2.290933505	1.48E-07	0.0089	94417.61756
		200 - 500	450.931	4.51E-07		34.60	4.53	39.13	6.26E-18	1.79E+10	1.12E-07	0.0067	124937.6456	
		500 - 1000	966.281	9.66E-07		80.61	20.79	101.41	1.62E-17	7.15E+09	1.17	1.16E-07	0.0070	120647.4815
		1000-2500	2254.655	2.25E-06		204.86	113.21	318.08	5.09E-17	2.81E+09	1.07	1.43E-07	0.0086	97969.3398
2500 - 5000		4831.404	4.83E-06		471.32	519.85	991.17	1.59E-16	1.26E+09	1.03	2.00E-07	0.0120	69932.23786	
RH - 80%	First iteration	100 - 200	100.4319169	1.00E-07		2.75	0.20	2.94	4.71018E-19	1.5891E+11	2.656047252	7.34273E-08	0.004405641	107652.90
		200 - 500	351.5117093	3.52E-07		13.38	2.40	15.79	2.5294E-18	24703894008	1.473156358	6.24006E-08	0.003744033	126676.22
		500 - 1000	753.2393771	7.53E-07		33.67	11.04	44.71	7.15358E-18	9553667313	1.2208063	6.83429E-08	0.004100575	115661.83
		1000-2500	1757.558547	1.76E-06		91.59	60.10	151.68	2.42691E-17	3671253871	1.094631271	8.90928E-08	0.005345891	88718.61
		2500 - 5000	3766.196886	3.77E-06		221.41	275.95	497.36	7.95771E-17	1634259144	1.04416126	1.3005E-07	0.007802978	60781.92
	5000 - 10000	7532.393771	7.53E-06		488.61	1103.79	1592.40	2.54784E-16	7998499273	1.02208063	2.03789E-07	0.012227358	38788.43	
	Second iteration	100 - 200	100.4319169	1.00E-07		6.38	0.22	6.61	1.06E-18	1.56E+11	2.656047252	1.63E-07	0.0099	87971.33066
		200 - 500	351.5117093	3.52E-07		26.20	2.75	28.96	4.63E-18	2.47E+10	1.47	1.14E-07	0.0069	126676.2248
		500 - 1000	753.2393771	7.53E-07		61.19	12.64	73.83	1.18E-17	9.55E+09	1.22	1.13E-07	0.0068	128474.6413
		1000-2500	1757.558547	1.76E-06		155.85	68.79	224.65	3.59E-17	3.67E+09	1.09	1.32E-07	0.0079	109872.7077
2500 - 5000		3766.196886	3.77E-06		359.17	315.89	675.06	1.08E-16	1.63E+09	1.04	1.77E-07	0.0106	82137.63957	
5000 - 10000	7532.393771	7.53E-06		764.17	1263.57	2027.74	3.24E-16	8.00E+08	1.02	2.60E-07	0.0156	55870.89967		
RH - 80%	First iteration	100 - 200	100.4319169	1.00E-07		2.73	0.19	2.93	4.6879E-19	1.5896E+11	2.6632	7.33132E-08	0.00441079	107527.22
		200 - 500	268.254	2.68E-07		9.59	1.40	10.99	1.75781E-18	35590171733	1.620099394	6.25748E-08	0.003754491	126233.39
		500 - 1000	574.829	5.75E-07		24.34	6.43	30.77	4.92326E-18	13221616446	1.289338221	6.59934E-08	0.003905607	121435.67
		1000-2500	1341.268	1.34E-06		66.72	35.00	101.72	1.62751E-17	4939783473	1.124002064	8.0955E-08	0.004823727	98322.31
		2500 - 5000	2874.146	2.87E-06		162.16	160.71	322.86	5.16583E-17	2169595981	1.057867624	1.12078E-07	0.006724663	70528.45
	5000 - 10000	5748.292	5.75E-06		359.25	642.83	1002.08	1.6333E-16	1055127604	1.028933812	1.69172E-07	0.010150325	46725.59	
	Second iteration	100 - 200	100.4319169	1.00E-07		6.35	0.22	6.57	1.05E-18	1.57E+11	2.6632	1.65E-07	0.0099	91341.62161
		200 - 500	268.254	2.68E-07		19.36	1.60	20.96	3.35E-18	3.56E+10	1.62	1.20E-07	0.0072	126233.3866
		500 - 1000	574.829	5.75E-07		45.33	7.36	52.69	8.43E-18	1.32E+10	1.29	1.11E-07	0.0067	135316.4068
		1000-2500	1341.268	1.34E-06		115.76	40.06	155.82	2.49E-17	4.94E+09	1.12	1.23E-07	0.0074	122477.33
2500 - 5000		2874.146	2.87E-06		267.28	183.97	451.25	7.22E-17	2.17E+09	1.06	1.57E-07	0.0094	96292.80256	
5000 - 10000	5748.292	5.75E-06		569.53	735.88	1305.42	2.09E-16	1.06E+09	1.03	2.20E-07	0.0132	68443.82008		

Table F.3.a.1. Salt 25% calculation results for the different relative humidity conditions.

	Salt 25%					
	RH 95%		RH 90%		RH 90%	
	First iteration	Second iteration	First iteration	Second iteration	First iteration	Second iteration
Concentration (Ni)	1.29E+13	1.29E+13	1.28E+13	1.26E+13	1.28E+13	1.19E+13
Number of plates	2082.294093	2082.294093	2111.270414	2141.244022	2105.389776	2255.273446
Velocity (m/s)	0.120117575	0.120117575	0.11846823	0.11680911	0.118799285	0.110900477
t (s)	49.95105823	49.95105823	50.64648994	51.36585654	50.50353464	54.10256271
Ni/T New	6.5E+14	6.5E+14	6.5E+14	6.5E+14	6.5E+14	6.5E+14

Table F.3.a.2. Salt 25% updated Nit and Nit check calculations.

		SALT 75%												
RH - 95%		Stage (dp [=] nm)	Midpoint (dp [=] nm)	Midpoint (dp [=] m)	Diffusion Charging	Field Charging	Total Charges	Charge (C)	Particle Mobility (m ² /Vs)	Slip Correction	Electric Mobility (m ² /Vs)	Settling Velocity (V/m)	Ac (m)	
	100 - 200	128.837	1.29E-07		3.80	0.32	4.12	6.58923E-19	60117570972	1.89580183	3.96129E-08	0.002376771	199548.02	
	200 - 500	450.931	4.51E-07		18.14	3.96	22.10	3.53584E-18	1138163201	1.255737195	4.02436E-08	0.002414616	196420.47	
	500 - 1000	966.281	9.66E-07		45.30	18.16	63.46	1.01538E-17	4734522175	1.119340024	4.8076E-08	0.002884417	164248.40	
	1000-2500	2254.655	2.25E-06		122.41	98.90	221.31	3.50809E-17	1905481878	1.051147439	6.74072E-08	0.00404821	117157.97	
	2500 - 5000	4831.404	4.83E-06		294.58	454.12	748.70	1.19792E-16	8661375607	1.023868805	1.07356E-07	0.006253585	76184.85	
	5000 - 10000	9662.809	9.66E-06		647.93	1816.46	2464.39	3.94303E-16	428020855.4	1.011934402	1.6877E-07	0.010126184	46836.99	
Second iteration		Stage (dp [=] nm)	Midpoint (dp [=] nm)	Midpoint (dp [=] m)	Diffusion Charging	Field Charging	Total Charges	Charge (C)	Particle Mobility (m ² /Vs)	Slip Correction	Electric Mobility (m ² /Vs)	Settling Velocity (V/m)	Ac (m)	
	100 - 200	128.837	1.29E-07		8.47	0.37	8.84	1.41E-18	6.01E+10	1.89580183	8.50E-08	0.0051	164618.1986	
	200 - 500	450.931	4.51E-07		34.60	4.53	39.13	6.26E-18	1.14E+10	1.26	7.13E-08	0.0043	166240.474	
	500 - 1000	966.281	9.66E-07		80.61	20.79	101.41	1.62E-17	4.73E+09	1.12	7.68E-08	0.0046	182209.0094	
	1000-2500	2254.655	2.25E-06		204.87	113.21	318.08	5.09E-17	1.91E+09	1.05	9.70E-08	0.0058	144337.8236	
	2500 - 5000	4831.404	4.83E-06		471.33	519.85	991.18	1.59E-16	8.66E+08	1.02	1.37E-07	0.0082	101900.6224	
	5000 - 10000	9662.809	9.66E-06		1001.45	2079.40	3080.86	4.93E-16	4.28E+08	1.01	2.11E-07	0.0127	66340.52811	
First iteration		Stage (dp [=] nm)	Midpoint (dp [=] nm)	Midpoint (dp [=] m)	Diffusion Charging	Field Charging	Total Charges	Charge (C)	Particle Mobility (m ² /Vs)	Slip Correction	Electric Mobility (m ² /Vs)	Settling Velocity (V/m)	Ac (m)	
	100 - 200	144.8478891	1.45E-07		4.41	0.41	4.82	7.71074E-19	87423202929	2.148239032	6.74098E-08	0.004044585	117262.95	
	200 - 500	506.9676117	5.07E-07		20.91	5.00	25.91	4.14626E-18	15441751588	1.328068295	6.40256E-08	0.003841535	123461.07	
	500 - 1000	1086.359168	1.09E-06		52.04	22.96	75.00	1.19996E-17	625657964	1.153098538	7.50787E-08	0.004504724	105285.04	
	1000-2500	2534.838059	2.53E-06		140.22	125.00	265.23	4.24361E-17	2478026389	1.065613659	1.05158E-07	0.00630946	75169.66	
	2500 - 5000	5431.79584	5.43E-06		336.77	573.99	910.77	1.45723E-16	1118466604	1.00619708	1.62981E-07	0.00977886	48500.41	
	5000 - 10000	10863.59168	1.09E-05		739.61	2295.97	3035.58	4.85092E-16	55091115.2	1.015309854	2.67574E-07	0.016054542	29542.00	
Second iteration		Stage (dp [=] nm)	Midpoint (dp [=] nm)	Midpoint (dp [=] m)	Diffusion Charging	Field Charging	Total Charges	Charge (C)	Particle Mobility (m ² /Vs)	Slip Correction	Electric Mobility (m ² /Vs)	Settling Velocity (V/m)	Ac (m)	
	100 - 200	144.8478891	1.45E-07		9.67	0.47	10.14	1.62E-18	8.74E+10	2.148239032	1.42E-07	0.0085	97108.48587	
	200 - 500	506.9676117	5.07E-07		39.42	5.72	45.15	7.22E-18	1.54E+10	1.33	1.12E-07	0.0067	123461.0651	
	500 - 1000	1086.359168	1.09E-06		91.75	26.28	118.03	1.89E-17	6.26E+09	1.15	1.18E-07	0.0071	116551.3509	
	1000-2500	2534.838059	2.53E-06		232.93	143.10	376.03	6.02E-17	2.48E+09	1.07	1.49E-07	0.0089	92370.44608	
	2500 - 5000	5431.79584	5.43E-06		535.47	657.08	1192.56	1.91E-16	1.12E+09	1.03	2.13E-07	0.0128	64330.80897	
	5000 - 10000	10863.59168	1.09E-05		1137.05	2628.32	3765.38	6.02E-16	5.51E+08	1.02	3.32E-07	0.0199	41492.2527	
First iteration		Stage (dp [=] nm)	Midpoint (dp [=] nm)	Midpoint (dp [=] m)	Diffusion Charging	Field Charging	Total Charges	Charge (C)	Particle Mobility (m ² /Vs)	Slip Correction	Electric Mobility (m ² /Vs)	Settling Velocity (V/m)	Ac (m)	
	100 - 200	110.5396	1.11E-07		3.11	0.24	3.35	5.36159E-19	1.33561E+11	2	5.04619159	1.16E-08	0.004296597	138585.02
	200 - 500	386.8887	3.87E-07		17.96	3.33	21.30	2.87422E-18	21783780254	1.428901073	6.23E-08	0.002421	126327.37	
	500 - 1000	829.0472	8.29E-07		37.75	13.37	51.13	1.8017E-18	853656222	1.200615839	6.98304E-08	0.004189817	113198.27	
	1000-2500	1934.4435	1.93E-06		102.43	72.80	175.23	2.80366E-17	3309187928	1.085978215	9.27785E-08	0.00556671	81599.33	
	2500 - 5000	4145.2626	4.15E-06		247.18	334.29	581.46	9.30444E-17	1479080721	1.040123168	1.37605E-07	0.00825624	57146.07	
	5000 - 10000	8290.4721	8.29E-06		544.76	1337.15	1881.91	3.0105E-16	725276318.7	1.020061584	2.18385E-07	0.013103081	36496.04	
Second iteration		Stage (dp [=] nm)	Midpoint (dp [=] nm)	Midpoint (dp [=] m)	Diffusion Charging	Field Charging	Total Charges	Charge (C)	Particle Mobility (m ² /Vs)	Slip Correction	Electric Mobility (m ² /Vs)	Settling Velocity (V/m)	Ac (m)	
	100 - 200	110.5396	1.11E-07		7.12	0.27	7.39	1.18E-18	1.34E+11	2.504619159	1.58E-08	0.0095	90554.22353	
	200 - 500	386.8887	3.87E-07		29.17	5.33	34.50	5.37E-18	2.18E+10	1.11E-07	1.06E-08	126327.37		
	500 - 1000	829.0472	8.29E-07		68.05	15.31	83.36	3.33E-18	3.54E+09	1.14E-07	0.0098	125615.8462		
	1000-2500	1934.4435	1.93E-06		173.17	83.34	256.51	4.10E-17	3.31E+09	1.09	1.36E-07	0.0081	105303.7804	
	2500 - 5000	4145.2626	4.15E-06		398.81	382.68	781.48	1.25E-16	1.48E+09	1.04	1.85E-07	0.0111	73730.5971	
	5000 - 10000	8290.4721	8.29E-06		848.06	1530.70	2378.76	3.81E-16	7.25E-08	1.02	2.76E-07	0.0166	51809.3186	

	Salt 25%					
	RH 95%		RH 90%		RH 90%	
	First iteration	Second iteration	First iteration	Second iteration	First iteration	Second iteration
Concentration (Ni)	8.22E+12	8.22E+12	1.31E+13	1.31E+13	1.28E+13	1.28E+13
Number of plates	3273.674566	3273.674566	2057.684418	2057.684418	2103.966	2103.966
Velocity (m/s)	0.076390119	0.076390119	0.121554867	0.121554867	0.118879716	0.118879716
t (s)	78.54418959	78.54418959	49.36042603	49.36042603	50.471184	50.471184
Ni _t New	6.5E+14	6.5E+14	6.5E+14	6.5E+14	6.5E+14	6.5E+14

Table F.3.a.2. Salt 25% updated Nit and Nit check calculations.

From these data, the following trends can be observed. First and foremost, we can observe how – across the different salt types and humidity conditions – the peak area (our limiting factor when it comes to achieving the desired efficiency) again occurs at the accumulation mode size. This size of particles will eventually determine the total number of plates required, and thus the total cost of the project. The graphs below illustrate this condition:

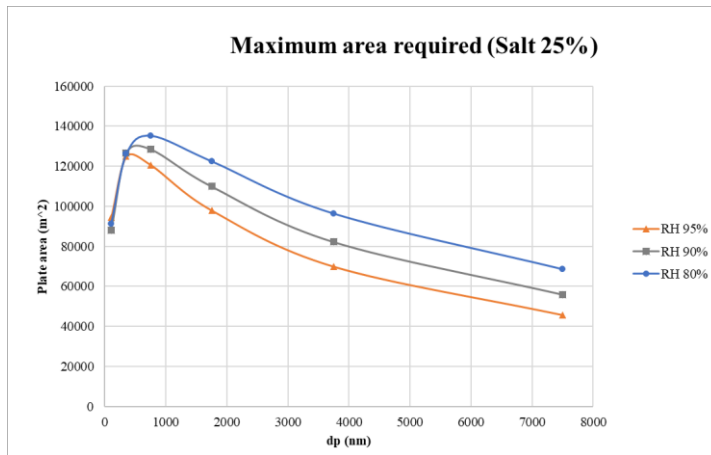


Figure F.1.a. Peak area occurring at accumulation mode for salt 25%.

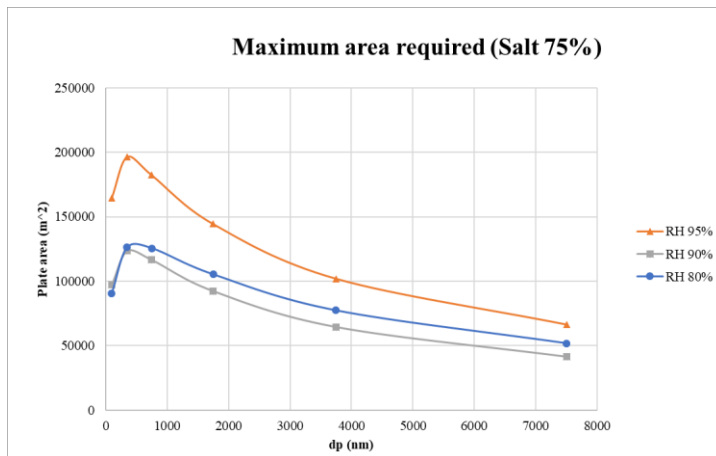


Figure F.1.a. Peak area occurring at accumulation mode for salt 25%.

Overall, the updated efficiency with the particle new concentration (Nit) appears to remain consistent across the different types of particles at different humidity values. A discrepancy occurs at Salt 75% in the 95% RH category, and this is because the particle growth and thus the plate are significantly higher compared to the other particles, which leads to a decrease in efficiency. This phenomenon can be appreciated in Figure F.1.a., with a peak close to 200,000 m², compared to an average of (around) 120,000 m². The table below summarizes the findings:

Updated Efficiency					
RH 95%		RH 90%		RH 80%	
Salt 25%	Salt 75%	Salt 25%	Salt 75%	Salt 25%	Salt 75%
0.97	0.90	0.97	0.96	0.97	0.97

Table F.4. Updated efficiencies with new Nit concentration.

Last but not least, the operation cost and the production cost can be observed in the tables below:

Capital Cost					
RH 95%		RH 90%		RH 80%	
Salt 25%	Salt 75%	Salt 25%	Salt 75%	Salt 25%	Salt 75%
\$ 697,154.29	\$927,085.46	\$ 709,523.99	\$ 691,952.09	\$ 733,099.47	\$ 701,716.67

Table F.5. Capital cost of production given by Cooper and Alley.

Operation Cost	
Daily	Yearly
\$80.40	\$29,344.32

Table F.5. Annual and Daily operational costs.

As I can be seen in Table F.4., the capital cost is in the range of ~\$700,000 on average, though there is spike in the cost of an ESP that would capture Salt 75% at 95% humidity because of the increased plate area. When compared to the operation costs reported in Table F.5., it can be seen that the main cost is the capital cost, which is almost twenty-five times greater. Should the device be operated for five years, and neglecting factors such as the time-value of money, interest rates, and inflation, among others, then the net cost of year operation would be – roughly - \$145,000/year (considering an average capital cost of ~\$700,000).