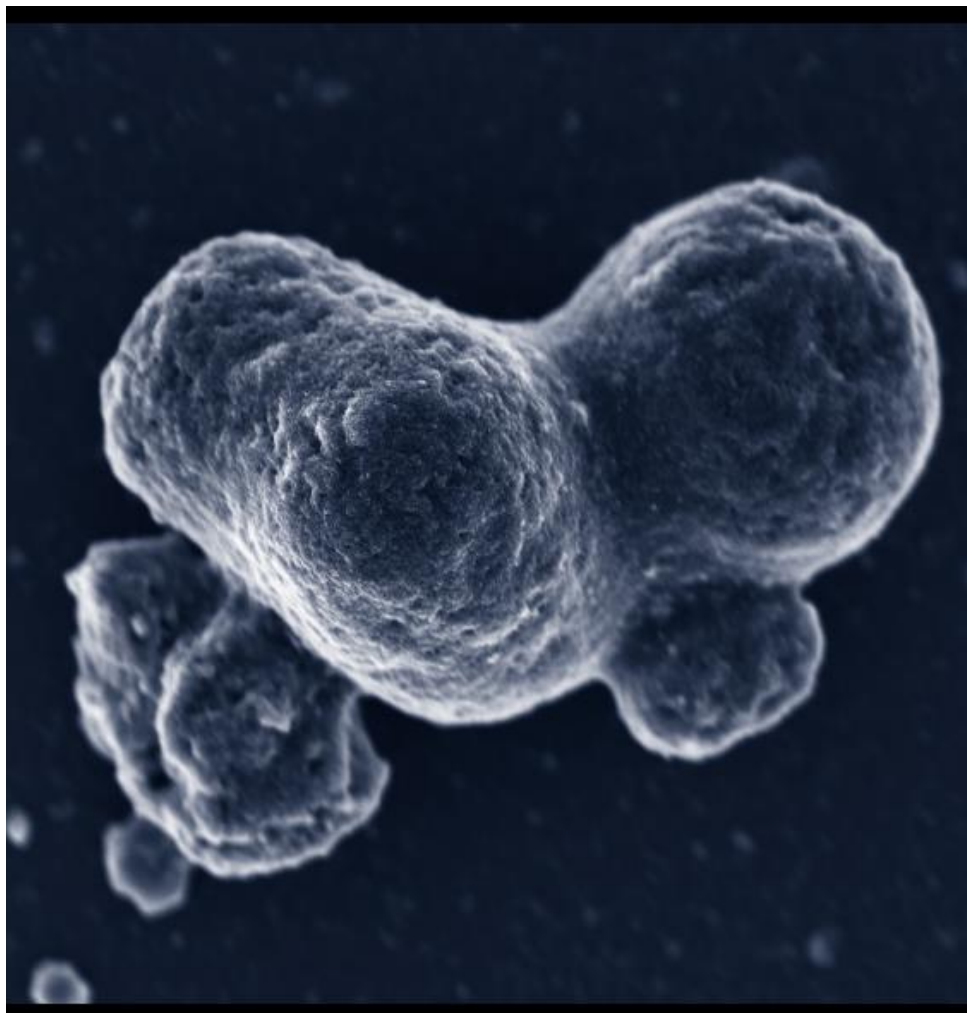




Reliable measurements of exposure to airborne engineered nanomaterials

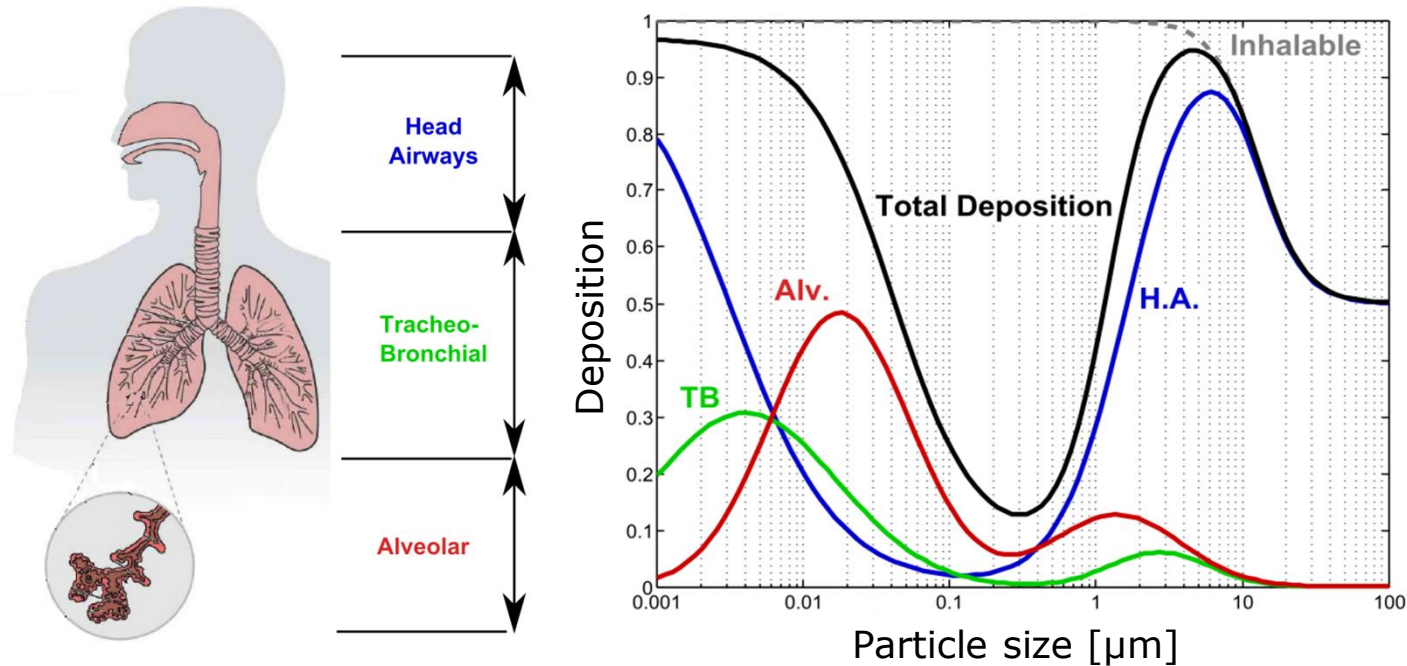
Marcus Levin

2016-01-11

Exposure to Engineered nanomaterials

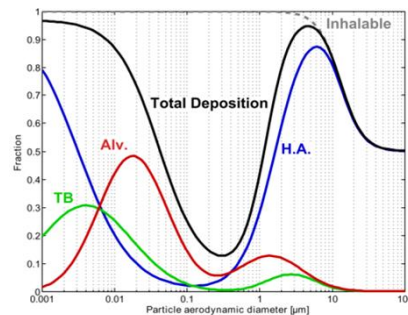
INHALATION

- Ç Most important route due to high uptake and strong toxic effects
- Ç Inhaled particle will deposit in different regions of the lungs; Head airways (**H.A.**), Tracheo-bronchial (**TB**) and Alveolar (**Alv.**) depending on particle size.
- Ç Unclear what metric drives toxic response – **Number, Mass, Surface Area**



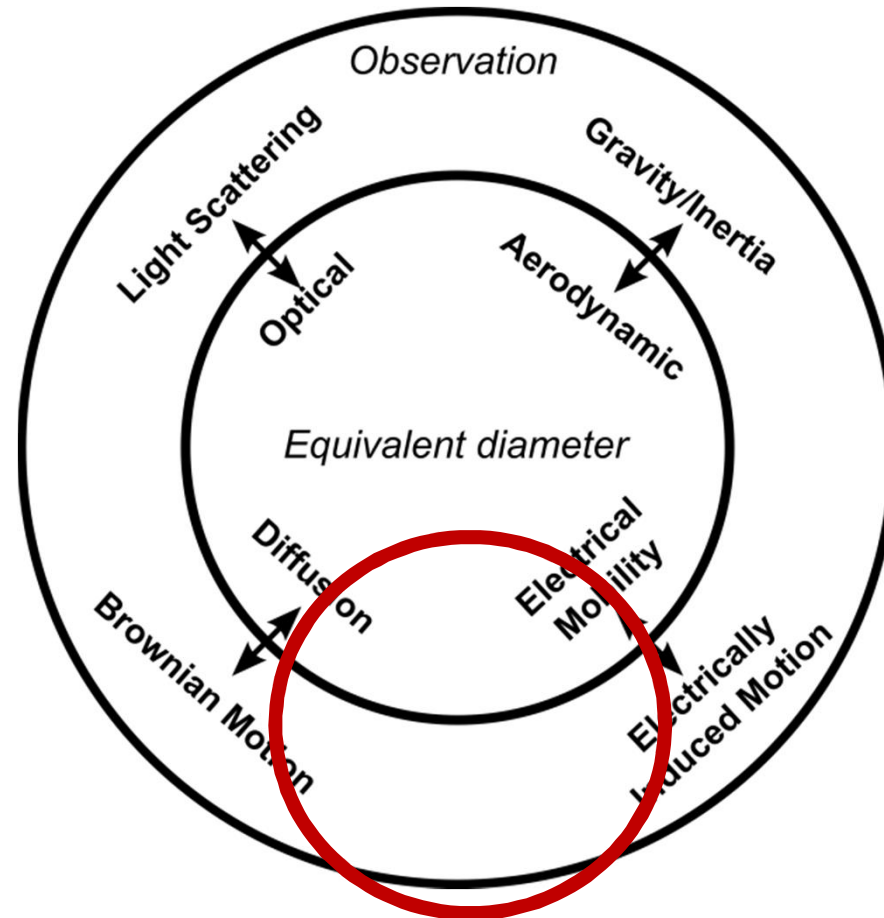
Measurement metrics

<i>Aerosol A</i>	High/Low	<i>Aerosol B</i>
Dp = 20 nm	Size	Dp = 200 nm
10 000 cm⁻³	Number	100 cm⁻³
12 μm²/cm³	Surface Area ~ D_p²	12 μm²/cm³
0.04 μm³/cm³	Volume ~ D_p³	0.4 μm³/cm³
6 μm²/cm³	Lung Deposited Surface Area (Alv)	0.8 μm²/cm³



Size is always important!

Size measurement

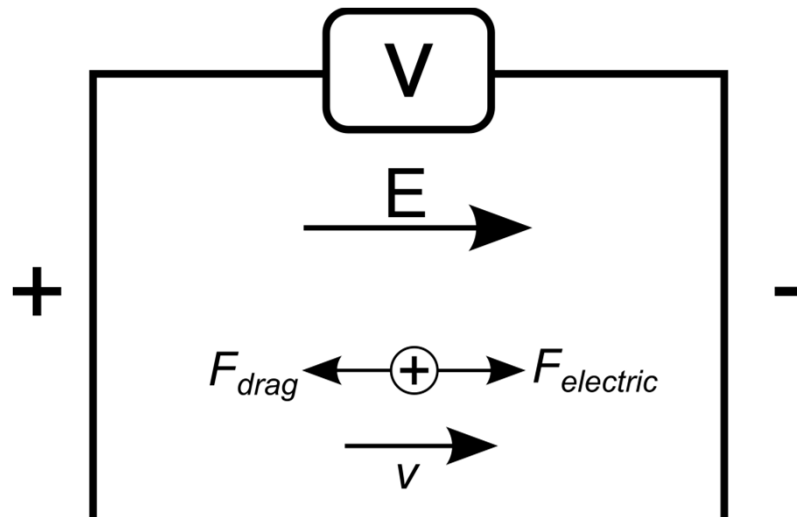


- **"True" particle size not feasible to measure**
- **Measured size is based on method of observation**

Electrical Mobility Measurements

Electrical Mobility, Z – Charged particles move through a medium in response to an electric field

Mobility equivalent diameter – Diameter of a sphere having the same mobility as the observed particle



Used in e.g.:

- Ç Scanning Mobility Particle Sizer
- Ç Fast Mobility Particle Sizer
- Ç NanoScan

With balance of electrostatic & drag forces: $\frac{q}{6\pi\eta r}$

Aerosol charging

- Objective is to create a known charge of each particle

- Diffusion Charging



(Unipolar or Bipolar)

$$Q_p \propto \frac{Q_p^2}{r_p}$$

$$Q_p = \frac{Q_p^2 r_p}{Q_p^2 r_p}$$

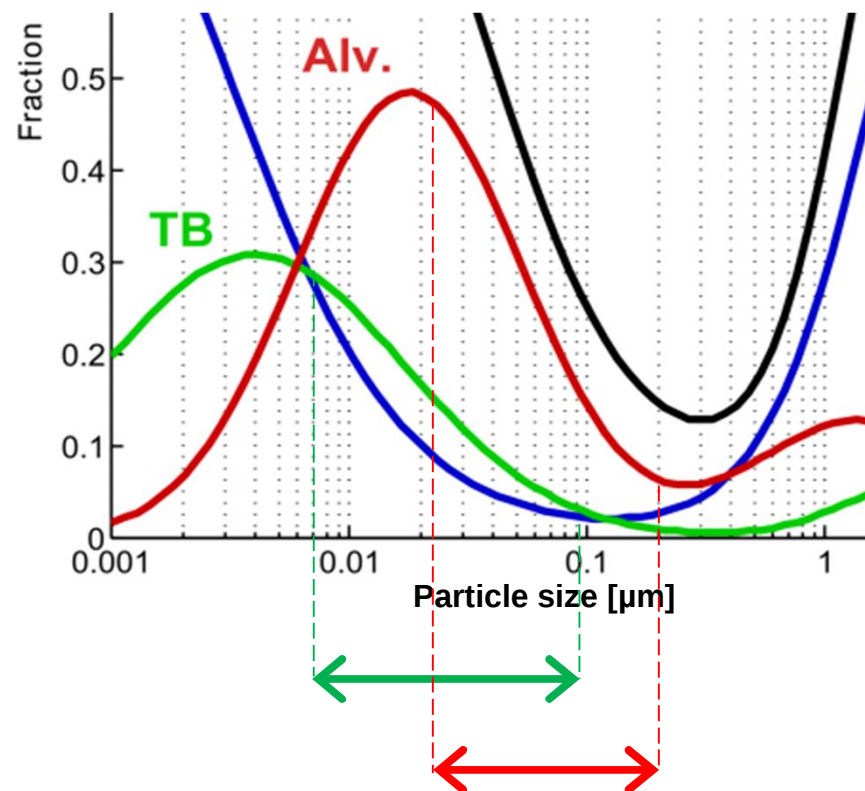
Lung Deposited Surface Area (LDSA)

Measurement of LDSA concentration:

- Deposition fractions $\sim \frac{m^2}{m^3}$
- Surface area $\sim \frac{m^2}{m^3}$
- **DF** ($\frac{m^2}{m^3}$) **X** **SA** ($\frac{m^2}{m^3}$) = **LDSA** ($\frac{m^2}{m^3}$)
- Diffusion charging with an electrometer yields a $\frac{m^2}{m^3}$ response



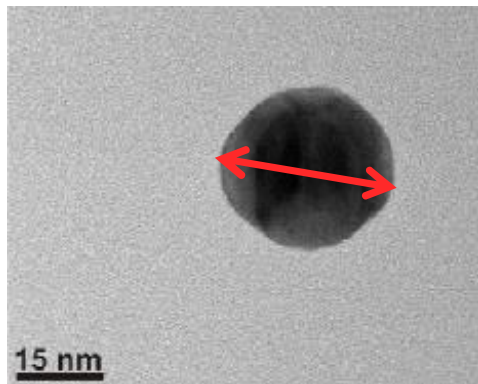
Nanoparticle Surface Area Monitor (NSAM)



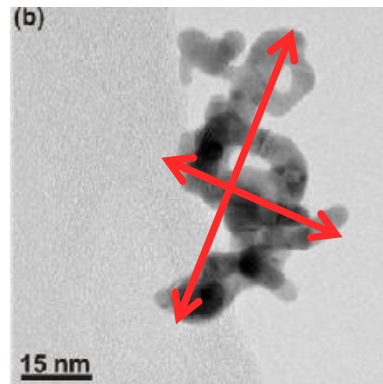
Non-spherical particles

Airborne particles are not just spheres with various diameters

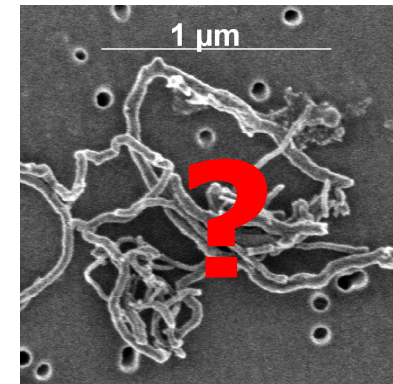
Spherical



Agglomerate



Fibrous



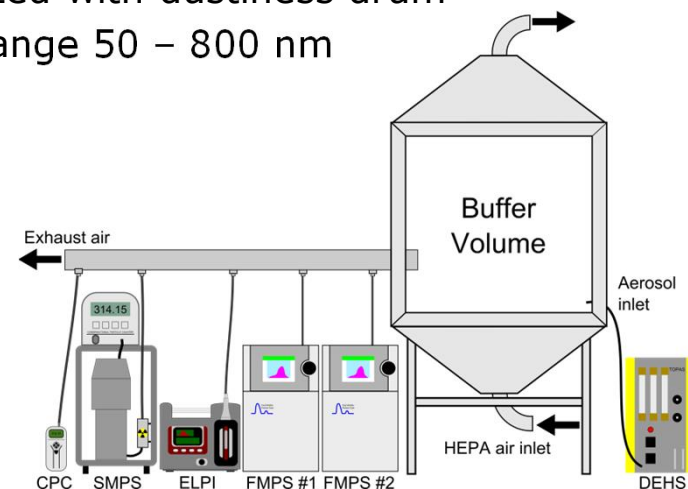
- **Charging effects from particle shape hard to predict**
- **Additional density/drag/etc. effects**
- **Agglomerate "problem" from powder handling**

Instrument reliability

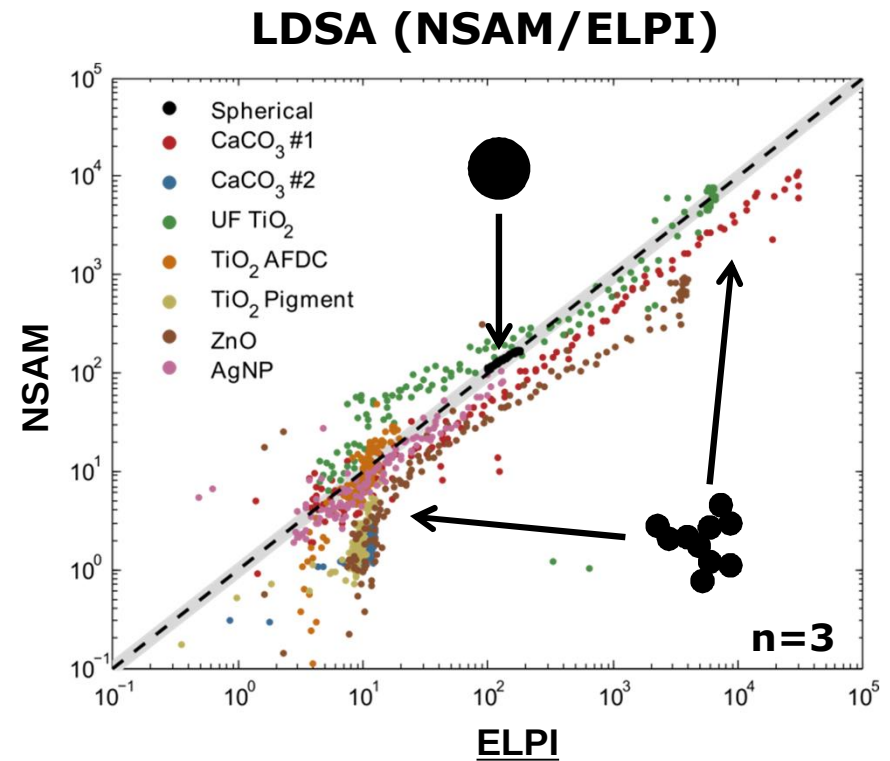
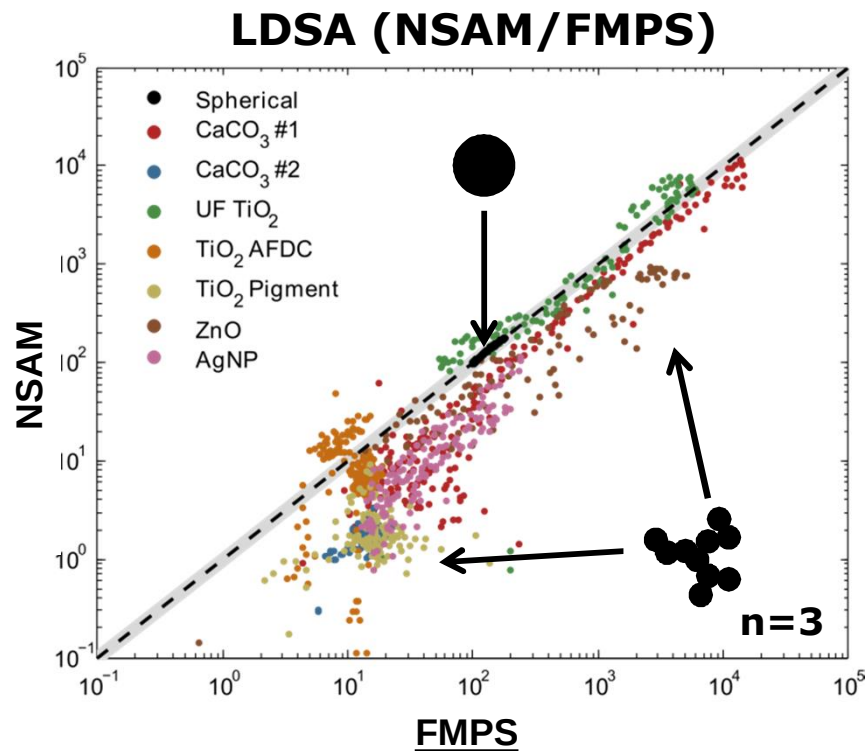
- Compare methods for real-time measurement of lung-deposited surface area (LDSA)
- Investigate to applicability of SMPS, FMPS, ELPI and NSAM in measurements of ENMs
 - Scanning Mobility Particle Sizer (SMPS)
 - Fast Mobility Particle Sizer (FMPS)
 - Electrical Low Pressure Impactor (ELPI)
 - Nanoparticle Surface Area Monitor

Test aerosols used:

- **LDSA measurements:** 7 powder nanomaterials aerosolized with dustiness drum
- **Instrument performance:** Spherical oil particles, size range 50 – 800 nm



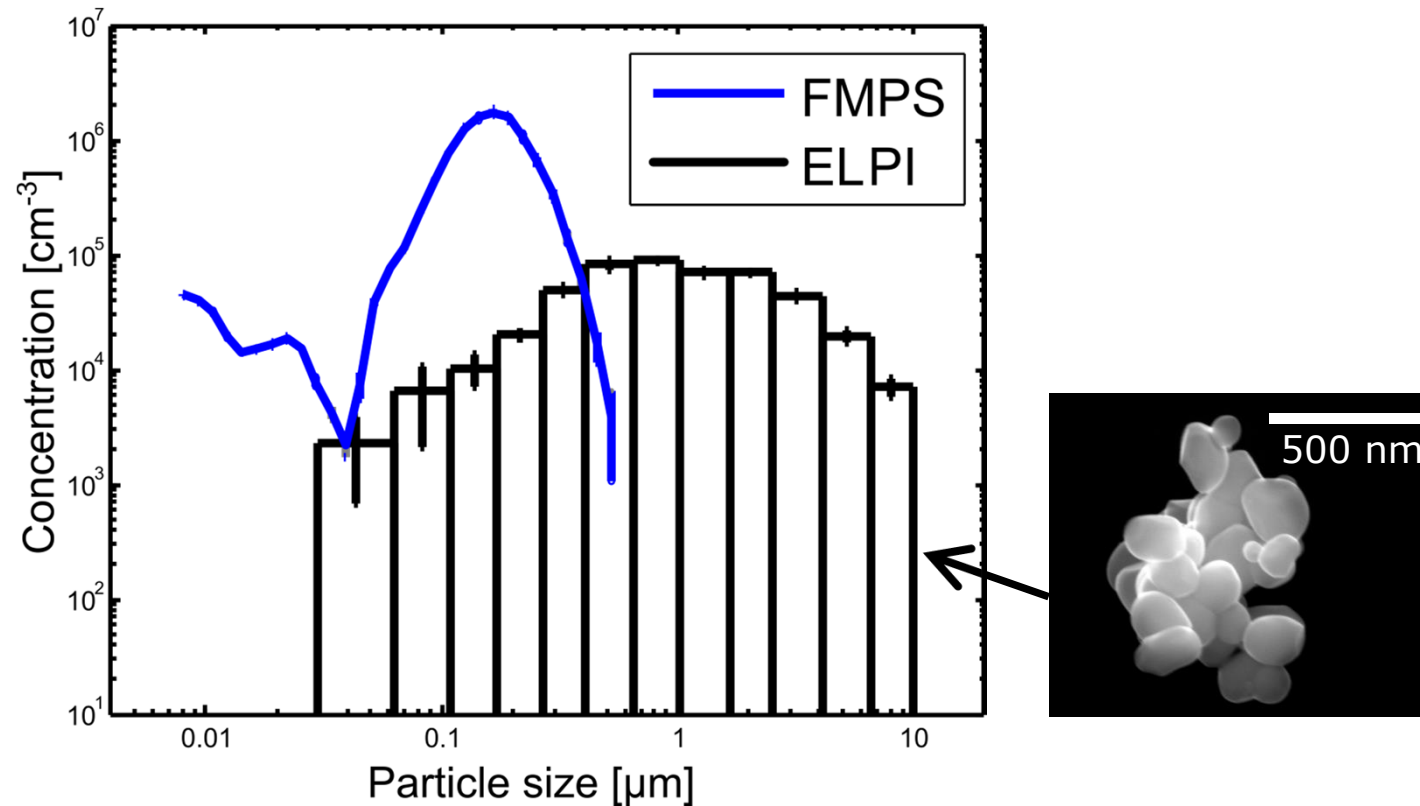
Measurement of LDSA



- **LDSA time-series comparison from dustiness generation**
- **Excellent agreement for spherical particles (40 nm)**
- **Material dependent for agglomerates**

Levin et al. 2015b

Measurements of complex particles



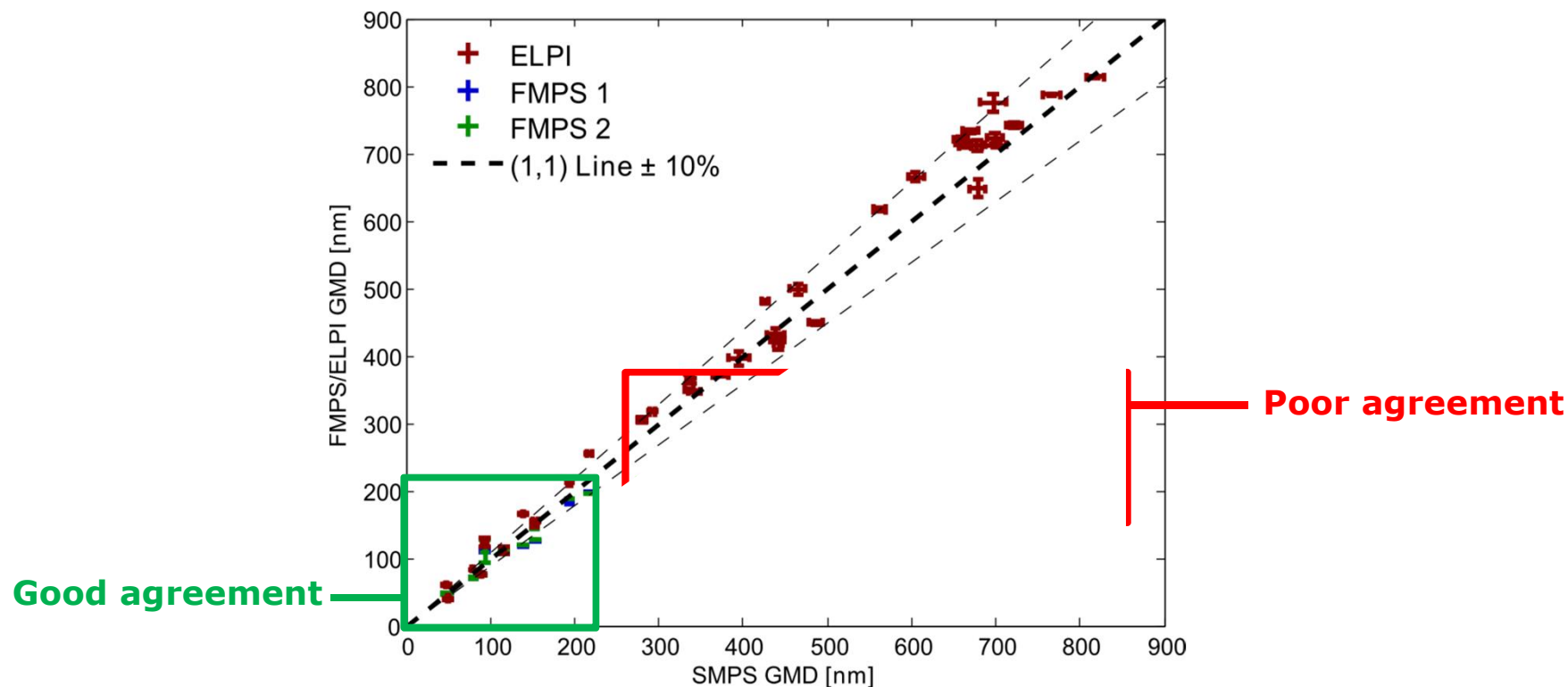
Compared to ELPI:

- **FMPS underestimate particle geometric mean diameter**
- **FMPS overestimate number concentration**

Levin et al. 2015b

Spherical particles

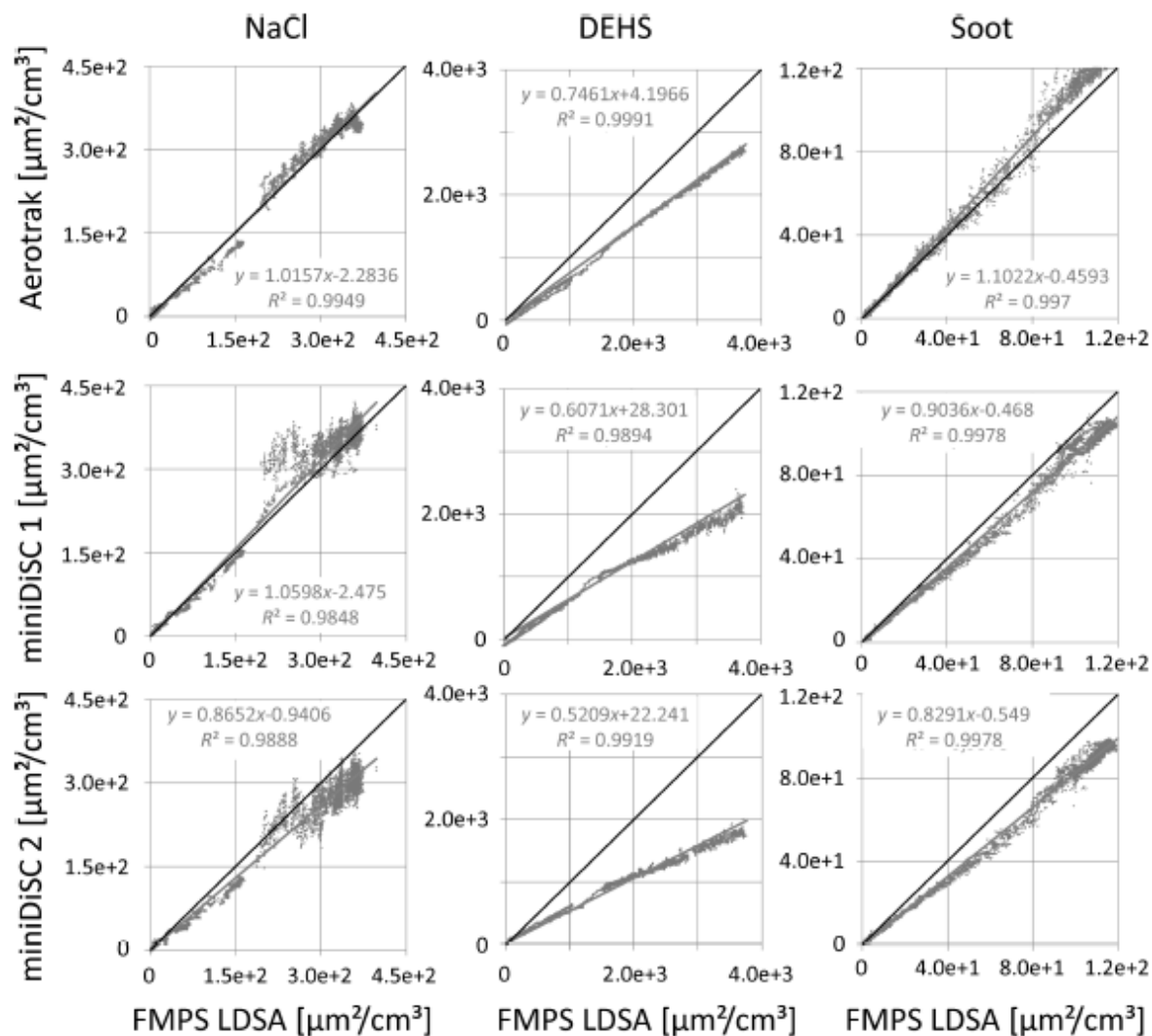
–Geometric Mean Diameter



- **Good agreement in GMD between SMPS and ELPI ($p=0.97$)**
- **Good agreement between SMPS and FMPS up until 200 nm**
- **GMD of 200-300 nm remains with FMPS as particles grow.**

Levin et al. 2015a

Comparison - LDSA



Correlation dependent on particle type!

Asbach et al. 2012

Key conclusions

- Current instrumentation is not reliable:
 - Specific instrument problems (e.g. FMPS)
 - General problem of non-sphericity
- Further development of strategies and techniques needed for occupational exposure limits to be based on lung deposited surface area
- Comparability can be achieved through:
 - Harmonization and "relative" measurements
 - New techniques for "absolute" measurements