

Supporting Information

We note that in order to obtain the sub-cooled liquid vapor pressures used in Figure 9, the solid vapor pressure of a compound, found in the literature, can be converted to that of the sub-cooled liquid from the following equation (39).

$$\ln \frac{P_s^\circ}{P_L^\circ} = -\frac{\Delta S_{\text{fus}}(T_M)}{R} \left[\frac{T_M}{T} - 1 \right] \quad (11)$$

where P_s° is the solid vapor pressure, P_L° sub-cooled liquid vapor pressure, T_m is the melting point, T is the temperature of interest and $\Delta_{\text{fus}}S$ is the entropy of fusion.

The derivation of equation (8) from equation (6), when $P \ll P^\circ$ and $C_{\text{BET}} \gg 1$, is as follows

$$\frac{n}{n_m} = \frac{C_{\text{BET}} x}{(1-x)(1+(C_{\text{BET}}-1)x)}, x = \frac{P}{P^\circ} \quad (6)$$

Equation (6) simplifies to the following

$$\frac{n}{n_m} \cong \frac{C_{\text{BET}} x}{1 + C_{\text{BET}} x} \quad (12)$$

since $1-x \approx 1$ and $C_{\text{BET}} - 1 \approx C_{\text{BET}}$. Equation (12) can be reformulated as follows

$$\frac{n}{n_m} \cong \frac{C_{\text{BET}} \frac{P}{P^\circ}}{1 + C_{\text{BET}} \frac{P}{P^\circ}} \times \frac{P^\circ}{P^\circ} \cong \frac{C_{\text{BET}} P}{P^\circ + C_{\text{BET}} P} \quad (13)$$

and since $\frac{n}{n_m} = \frac{\theta}{\theta_m}$ we can equate (13) to the Langmuir isotherm (1) as follows

$$\frac{C_{\text{BET}} P}{P^\circ + C_{\text{BET}} P} \cong \frac{KP}{1 + KP} \quad (14)$$

which simplifies to

$$K \cong \frac{C_{\text{BET}}}{P^{\circ}} \quad (8)$$

- (39) Schwarzenbach, R. P.; Gschwend, P. M.; Imboden, D. M. *Environmental Organic Chemistry*; 2 ed.; Wiley-Interscience: Hoboken, 2003.